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# UTILITY PATENT APPLICATION TRANSMITTAL

(Only for new nonprovisional applications under 37 CFR 1.53(b))

Attorney Docket No.	101900
First Inventor	David G. Boyers et al.
Title	Method and Apparatus for Treating a Substrate With an Ozone-Solvent Solution
Express Mail Label No.	EK 798 663 677 US

## APPLICATION ELEMENTS

See MPEP chapter 600 concerning utility patent application contents.

## ADDRESS TO:

 Assistant Commissioner for Patents  
 Box Patent Application  
 Washington, DC 20231

1. ☒ Fee Transmittal Form (e.g., PTO/SB/17)  
(Submit an original, and a duplicate for fee processing)
2. ☒ Applicant claims small entity status.  
See 37 CFR 1.27.
3. ☒ Specification [Total Pages 95]  
(preferred arrangement set forth below)  
 - Descriptive title of the Invention  
 - Cross References to Related Applications  
 - Statement Regarding Fed sponsored R & D  
 - Reference to sequence listing, a table, or a computer program listing appendix  
 - Background of the Invention  
 - Brief Summary of the Invention  
 - Brief Description of the Drawings (if filed)  
 - Detailed Description  
 - Claim(s)  
 - Abstract of the Disclosure
4. ☒ Drawing(s) (35 U.S.C. 113) [Total Sheets 19]
5. Oath or Declaration [Total Pages 2]
- a. ☒ Newly executed (original or copy)
- b. ☐ Copy from a prior application (37 C.F.R. § 1.63(d))  
(for continuation/divisional with Box 17 completed)
- i. ☐ **DELETION OF INVENTOR(S)**  
Signed statement attached deleting inventor(s) named in the prior application, see 37 C.F.R. §§ 1.63(d)(2) and 1.33(b).
6. ☐ Application Data Sheet. See 37 CFR 1.76.

7. ☐ CD-ROM or CD-R in duplicate, large table or Computer Program (Appendix)
8. Nucleotide and/or Amino Acid Sequence Submission (if applicable, all necessary)
- a. ☐ Computer Readable Copy (CRF)
- b. Specification Sequence Listing on:  
 i. ☐ CD-ROM or CD-R (2 copies); or  
 ii. ☐ paper
- c. ☐ Statement verifying identity of above copies

## ACCOMPANYING APPLICATION PARTS

9. ☐ Assignment Papers (cover sheet & document(s))
10. ☐ 37 C.F.R. § 3.73(b) Statement ☒ Power of Attorney  
(when there is an assignee)
11. ☐ English Translation Document (if applicable)
12. ☐ Information Disclosure Statement (IDS)/PTO-1449 ☐ Copies of IDS Citations
13. ☐ Preliminary Amendment
14. ☒ Return Receipt Postcard (MPEP 503)  
(Should be specifically itemized)
15. ☐ Certified copy of Priority Document(s) (if foreign priority is claimed)
16. ☒ Other: small entity Verification
- Other:

17. If a CONTINUING APPLICATION, check appropriate box, and supply the requisite information below and in a preliminary amendment, or in an Application Data Sheet under 37 CFR 1.76.

☐ Continuation ☐ Divisional ☒ Claims priority to Provisional application No. 60 / 160,435 filed October 19, 1999

Prior application information:

Examiner:

Group Art Unit:

For CONTINUATION OR DIVISIONAL APPS only: The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 5b, is considered a part of the disclosure of the accompanying continuation or divisional application and is hereby incorporated by reference. The incorporation can only be relied upon when a portion has been inadvertently omitted from the submitted application parts.

## 18. CORRESPONDENCE ADDRESS

☐ Customer Number or Bar Code Label (Insert Customer No. or Attach bar code label here) or ☒ Correspondence address below

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Name (Print/Type)	Joseph H. Smith	Registration No. (Attorney/Agent)	30,328
Signature		Date	October 19, 2000

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# FEE TRANSMITTAL for FY 2001

Patent fees are subject to annual revision.

## Complete if Known

Application Number	To be assigned
Filing Date	October 19, 2000
First Named Inventor	David G. Boyers et al
Examiner Name	To be assigned
Group Art Unit	To be assigned
Attorney Docket No	101900

TOTAL AMOUNT OF PAYMENT (\$)**1701**

## METHOD OF PAYMENT (check one)

1. ☒ The Commissioner is hereby authorized to charge indicated fees and credit any overpayment to:

Deposit Account Number: **10-1318**  
Account Name: **Joseph H. Smith**

- ☒ Charge any Additional Fee Required Under 37 CFR 1.16 and 1.17  
☒ Applicant claims small entity status. See 37 CFR 1.27

2. ☒ Payment Enclosed:  
☒ Check ☐ Credit card ☐ Money ☐ Other Order

## FEE CALCULATION

### 1. BASIC FILING FEE

Large Entity	Small Entity	Fee Code (\$)	Fee Code (\$)	Fee Description	Fee Paid
101	710	201	355	Utility filing fee	355
106	320	206	160	Design filing fee	
107	490	207	245	Plant filing fee	
108	710	208	355	Reissue filing fee	
114	150	214	75	Provisional filing fee	
SUBTOTAL (1)					(\$) <b>355</b>

### 2. EXTRA CLAIM FEES

Total Claims	Extra Claims	Fee from Allow.	Fee Paid
<b>119</b>	<b>99</b>	<b>9</b>	<b>891</b>
Independent Claims	<b>11</b>	<b>40</b>	<b>320</b>
Multiple Dependent	<b>105</b>	<b>135</b>	

\*or number previously paid, if greater. For Reissues, see below

Large Entity	Small Entity	Fee Code (\$)	Fee Code (\$)	Fee Description
103	18	203	9	Claims in excess of 20
102	80	202	40	Independent claims in excess of 3
104	270	204	135	Multiple dependent claim, if not paid
109	80	209	40	** Reissue independent claims over original patent
110	18	210	9	** Reissue claims in excess of 20 and over original patent

SUBTOTAL (2)

(\$)**1355**

## FEE CALCULATION (continued)

### 3. ADDITIONAL FEES

Large Entity	Small Entity	Fee Code (\$)	Fee Code (\$)	Fee Description	Fee Paid
105	130	205	65	Surcharge - late filing fee or oath	
127	50	227	25	Surcharge - late provisional filing fee or cover sheet.	
139	130	139	130	Non-English specification	
147	2520	147	2520	For filing a request for ex parte reexamination	
112	920*	112	920*	Requesting publication of SIR prior to Examiner action	
113	1840*	113	1840*	Requesting publication of SIR after Examiner action	
115	110	215	55	Extension for reply within first month	
116	390	216	195	Extension for reply within second month	
117	890	217	445	Extension for reply within third month	
118	1390	218	695	Extension for reply within fourth month	
128	1890	228	945	Extension for reply within fifth month	
119	310	219	155	Notice of Appeal	
120	310	220	155	Filing a brief in support of an appeal	
121	270	221	135	Request for oral hearing	
138	1510	138	1510	Petition to institute a public use proceeding	
140	110	240	55	Petition to revive - unavoidable	
141	1240	241	620	Petition to revive - unintentional	
142	1240	242	620	Utility issue fee (or reissue)	
143	440	243	220	Design issue fee	
144	600	244	300	Plant issue fee	
122	130	122	130	Petitions to the Commissioner	
123	50	123	50	Petitions related to provisional applications	
126	240	126	240	Submission of Information Disclosure Stmt	
581	40	581	40	Recording each patent assignment per property (times number of properties)	
146	710	246	355	Filing a submission after final rejection (37 CFR 1.129(a))	
149	710	249	355	For each additional invention to be examined (37 CFR 1.129(b))	
179	710	279	355	Request for Continued Examination (RCE)	
Other fee (specify) _____					
Other fee (specify) _____					

\* Reduced by Basic Filing Fee Paid

SUBTOTAL (3)

(\$)

## SUBMITTED BY

Name (Print/Type)	Registration No. (Attorney/Agent)	Telephone
Joseph H. Smith	30,328	408-270-3089
Signature	Date	
<i>Joseph H. Smith</i>	Oct. 19, 2000	

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**STATEMENT CLAIMING SMALL ENTITY STATUS  
(37 CFR 1.9(f) & 1.27(c))--SMALL BUSINESS CONCERN**

Docket Number (Optional)

Applicant, Patentee, or Identifier: Phifer Smith Corporation

Application or Patent No.: \_\_\_\_\_

Filed or Issued: \_\_\_\_\_

Title: METHOD AND APPARATUS FOR TREATING A SUBSTRATE WITH AN  
OZONE-SOLVENT SOLUTION

I hereby state that I am

☐ the owner of the small business concern identified below.

☒ an official of the small business concern empowered to act on behalf of the concern identified below

NAME OF SMALL BUSINESS CONCERN Phifer Smith Corporation

ADDRESS OF SMALL BUSINESS CONCERN 2181 Park Blvd.  
Palo Alto, CA 94306

I hereby state that the above identified small business concern qualifies as a small business concern as defined in 13 CFR Part 121 for purposes of paying reduced fees to the United States Patent and Trademark Office. Questions related to size standards for a small business concern may be directed to: Small Business Administration, Size Standards Staff, 409 Third Street, SW, Washington, DC 20416.

I hereby state that rights under contract or law have been conveyed to and remain with the small business concern identified above with regard to the invention described in:

☒ the specification filed herewith with title as listed above.

☐ the application identified above.

☐ the patent identified above.

If the rights held by the above identified small business concern are not exclusive, each individual, concern, or organization having rights in the invention must file separate statements as to their status as small entities, and no rights to the invention are held by any person, other than the inventor, who would not qualify as an independent inventor under 37 CFR 1.9(c) if that person made the invention, or by any concern which would not qualify as a small business concern under 37 CFR 1.9(d), or a nonprofit organization under 37 CFR 1.9(e).

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Separate statements are required from each named person, concern or organization having rights to the invention stating their status as small entities. (37 CFR 1.27)

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NAME OF PERSON SIGNING David G. Boyers

TITLE OF PERSON IF OTHER THAN OWNER President

ADDRESS OF PERSON SIGNING 2181 Park Blvd., Palo Alto, CA 94306

SIGNATURE David G. Boyers DATE Oct. 18, 2000

# A METHOD AND APPARATUS FOR TREATING A SUBSTRATE WITH AN OZONE-SOLVENT SOLUTION

## BACKGROUND - FIELD OF INVENTION

This invention presents a method for treating materials using an ozone-solvent solution. The method may be used for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like at high speed using a solution of ozone gas dissolved in a solvent. The method may also be used for disinfection or sterilization of medical instruments whereby bacteria, viruses, and other microbes are inactivated by the ozone-solvent solution.

## BACKGROUND - WAFER PROCESSING

Low Temperature Immersion Processing: The patent by Mathews (US Patent 5464480), publications by Kashkoush, et.al 1997, Kashkoush, et.al. 1998, and Mathews, 1998 disclose a method of removing photoresist from semiconductor wafers using with ozone dissolved in water at a temperature of 1 to 15 degree C and applied to the surface of the wafers at the same temperature. They achieved an etch rate for I-line positive photoresist of about 700 Angstroms/min at a dissolved concentration of about 90 mg/L and temperature of 5 degree C. They reported that DUV positive resists were not effectively removed by the process. The method has a number of disadvantages: 1) low etch rates for I-line positive photoresist and inability to etch DUV positive resists; this is a severe disadvantage because most high resolution lithography for electronic device fabrication is based upon positive photoresist with new generation designs using DUV resists, 2) the presence of high concentration ozone gas in the process tank presents a safety hazard to the users 3) the immersion design uses water inefficiently and provides no means for integration of a drying step. We can analyze the etch rate performance of this method by noting the following: 1) a low temperature of 1 to 15 degree C produces a higher dissolved ozone concentration C for a given gas phase concentration and pressure, 2) a low temperature of 1 to 15 degree C produces a low surface reaction rate, and 3) the very low mass transport available in an immersion design with a very large stagnant layer thickness  $\delta$  reduces the temperature at which the reaction becomes surface reaction rate limited, and 4) the etch rate is therefore lowest of the four prior art methods reviewed.

Room Temperature Spin Processing: The publications by Christenson, et. al. 1997, Christenson, et. al, 1998, and Nelson et. al. 1999 disclose a method of removing photoresist

from semiconductor wafers using ozone dissolved in water at room temperature (20 degreeC) and applied to the surface of the wafers at the same temperature. The method achieves improved mass transfer over that which can be achieved by an immersion process by applying an ozone-water solution to the center of a semiconductor wafer which is rotating at 1,000 rpm. The reported etch rate for I-line positive photoresist is non-uniform, peaking at 2500 Angstroms/min in the very center of the wafer and falling quickly to about 1500 Angstroms/minute at a radius of 30 mm and remaining relatively constant to the edge at a radius of 75 mm. No etch rate results were reported for DUV positive resists. The water is sent to waste after one pass over the wafer. The resist removal method offers some advantages over the method disclosed by Mathews and Kashkoush: 1) the reported etch rate for I-line positive photoresist is a factor of two higher (1500 Angstroms/min compared to 700 Angstroms/min), 2) high concentration ozone gas is not present in the process chamber, and 3) the use of a spinning wafer configuration provides for more efficient use of water and means for integration of a spin rinse and spin drying step at the end of the process. However, the photoresist etch rates which can be achieved with this technique are still too low to form the basis for a single wafer processing tool with a practical throughput. Our own measurements show that the etch rate for DUV positive photoresist that can be achieved with this technique is about 600 Angstroms per minute as compared to the 1500 Angstroms/min. for I-line positive photoresist. We can analyze the etch rate performance of this method by noting the following: 1) a temperature of 20 degree C produces an intermediate dissolved ozone concentration for a given gas phase concentration and pressure, 2) a temperature of 20 degree C produces a modest surface reaction rate, 3) the higher mass transport available in 1000 rpm spinning wafer design with a smaller stagnant layer thickness  $\delta$  increases the temperature at which the reaction becomes surface reaction rate limited, and 4) the etch rate is higher than that achieved by the low temperature immersion processing method.

Humidified Ozone Gas Processing Method: The publications by DeGendt, et. al. 1998 disclose a method of removing photoresist from semiconductor wafers using humidified ozone gas. In the moist gas phase process quartz container is filled with DI water. A diffuser is located in the bottom of the quartz container and high concentration ozone gas is bubbled through the liquid while the liquid is heated to about 80 degree C. One or more photoresist coated semiconductor wafers are placed above the liquid level (not immersed) and exposed to a moist ozone gas ambient. Etch rates as high as 6,000 Angstroms/min. for I-line positive and DUV positive photoresist were reported. These workers also noted that the ozone etch process

is limited by kinetic factors at low temperature and by ozone solubility at higher temperatures. They also note that any optimized process should aim at maximizing the ozone concentration at elevated temperatures. They state that by exposing the wafer to a moist gas ambient that a thin condensation layer is formed on the wafer and the ozone gas ambient maintains a continuous supply of ozone gas above the condensation layer and achieve a higher dissolved ozone concentration in the condensation layer. The method has several significant disadvantages: 1) the high concentration ozone gas in the presence of water vapor can be corrosive to metals, 2) the presence of high concentration ozone gas in the process tank presents a safety hazard to the users, and 3) the high etch rates can only be achieved at relatively high temperatures (80 C) where metal corrosion rates are increased further. Accordingly, the method may only be suitable for front end of line wafer processing before metals are deposited on the wafer. We can analyze the etch rate performance of this method by noting the following: 1) a temperature of 80 degree C produces very low dissolved ozone concentration for a given gas phase concentration and pressure, 2) a temperature of 80 degree C produces a high surface reaction rate, 3) the higher mass transport available in wet ozone gas method with a smaller stagnant layer thickness  $\delta$  increases the temperature at which the reaction becomes surface reaction rate limited, and 4) the etch rate is higher than that achieved by the low temperature immersion processing method or the room temperature spin processing method.

Humidified Ozone Gas with Spin Processing Method: Workers at Semitool (Scranton, 1999) have developed a similar wet ozone gas technique in which hot DI water at a temperature of between 40 and 95 deg. is sprayed onto the surface of a wafer spinning at 1,000 rpm in process chamber filled with 14 weight percent ozone gas. They have reported etch rates for I-line positive photoresist of 4200 angstroms/minute at a temperature of 95 deg. C, 2000 angstroms/minute at 45 deg. C, and 1200 angstroms/minute at 20 deg. C. They reported that in the case of back-end-of-line (BEOL) processes that metal corrosion limited process temperatures to about 45 deg. C. They reported that the 95 deg. C process was only suitable for use in front-end-of-line (FEOL) processes where metal corrosion was not a consideration. This method has the same disadvantages as the humidified ozone gas processing method: 1) the high concentration ozone gas in the presence of water vapor can be corrosive to metals, 2) the presence of high concentration ozone gas in the process tank presents a safety hazard to the users, and 3) the high etch rates can only be achieved at relatively high temperatures (95 C) where metal corrosion rates are increased further. We can analyze the etch rate

performance of this method by noting the following: 1) a temperature of 80 degree C produces very low dissolved ozone concentration for a given gas phase concentration and pressure, 2) a temperature of 80 degree C produces a high surface reaction rate, 3) the higher mass transport available in wet ozone gas method increases the temperature at which the reaction becomes surface reaction rate limited, and 4) the etch rate is higher than that achieved by the low temperature immersion processing method or the room temperature spin processing method.

Underlying Basis of the Etch Rate Limitations of the Prior Art Methods: In all the prior art described above, the ozone is dissolved in water at a given temperature and the ozone water solution is applied to the material to be oxidized at the same temperature. Accordingly, if one lowers the temperature to increase the dissolved ozone concentration, then one decreases the surface reaction. Alternatively, if one increases the temperature to increase the surface reaction rate, then one decreases the dissolved ozone concentration. Accordingly, the etch rate may be limited by a low dissolved ozone concentration or a low surface reaction rate.

#### BACKGROUND: MEDICAL INSTRUMENT STERILIZATION:

Humidified Ozone Gas Sterilization Method: Faddis, et. al discloses a technique for medical instrument sterilization with humidified ozone gas in US Patent 5,344,622 and US Patent 5,069,880. Karlson discloses a technique for medical instrument sterilization with humidified ozone gas in US Patent 5,069,880. This method has a number disadvantages including: 1) the high concentration ozone gas in the presence of water vapor can be corrosive to metals, 2) the high concentration ozone gas in the presence of water vapor is not compatible with many of the common plastic and elastomer materials used in the construction of medical devices, 3) the presence of high concentration ozone gas in the process tank presents a safety hazard to the users, and 4) the inactivation rate is relatively low because the surface reaction rate is relatively low.

Room Temperature Immersion Processing: Rickloff, 1987, and Omi, et. al A room temperature (20 deg. C) immersion processor has a number of limitations: 1) the modest temperature produces a modest surface reaction rate, 2) the very low mass transport available in an immersion design with a very large stagnant layer thickness  $\delta$  reduces the temperature at which the reaction becomes surface reaction rate limited, and 3) the inactivation rate is low because of the low surface reaction rate and low mass transport rate.

## SUMMARY OF INVENTION

In accordance with preferred embodiments of the invention, methods and apparatus are provided for treating materials which have broad advantages over prior systems and a very wide range of applications, including materials processing, wafer processing, medical instrument sterilization, and the like). The method entails forming an ozone-solvent solution at a first temperature having a dissolved ozone concentration and reacting the ozone-solvent solution with the material at a second temperature which is higher than said first temperature. In a preferred mode the reacting step comprises heating at least one of the ozone-solvent solution and the material, thereby causing said ozone-solvent solution to have a higher dissolved ozone concentration while reacting with the material than if the ozone-solvent solution had been formed at the second temperature. Various apparatus are presented to carry out these treatment processes.

For example, in a preferred embodiment of the invention, a system is provided for treating a substrate with an ozone-solvent solution which comprises a supply of an ozone-solvent solution formed at a first temperature which delivers a generally continuous supply of the ozone-solvent solution at the first temperature. The system includes a heater coupled to receive the ozone-solvent solution at the first temperature from the supply and heats the ozone-solvent solution received, and provides a generally continuous supply of heated ozone-solvent solution. The systems also includes an applicator fluidly coupled the heater to receive the generally continuous supply of the heated ozone-solvent solution, the applicator having an outlet configured to direct the heated ozone-solvent solution at a second temperature greater than the first temperature toward a substrate.

Outlines below are some of the advantages the various processes and how they are achieved:

higher oxidation rate: provides a method for oxidizing materials using a solution of ozone gas dissolved in solvent which can produce much higher oxidation rates than can be achieved by current methods

environmentally benign chemical: provides a method for oxidizing materials at high speed which uses an environmentally benign, residue free chemistry thereby reducing chemical disposal cost

increased user safety and reduced chemical cost and reduced chemical disposal cost: provides a method for oxidizing materials at high speed where the oxidizing chemical can be created and destroyed at the point of manufacture thereby increasing user safety, reducing chemical cost and reducing chemical disposal cost



additional chemicals can be injected with minimal impact on dissolved ozone concentration or injected-chemical concentration: provides a method for oxidizing materials using a solution of ozone gas dissolved in a solvent which may include injecting additional chemicals which significantly reduce the time available for injected chemicals to react with the ozone-solvent solution and thereby minimizes any decrease in dissolved ozone concentration or injected chemical concentration caused by such a reaction.

different injected chemicals may be added to the ozone-water solution during different phases of the materials processing cycle: provides a method for oxidizing materials using a solution of ozone gas dissolved in water which may include a means to select the mix of chemical(s) that are dispensed onto the surface of the materials to be oxidized during a given phase of the materials processing cycle

DI water and other chemicals may be introduced in lieu of the ozone-water solution during different phases of the materials processing cycle: provides a method for oxidizing materials using a solution of ozone gas dissolved in water which may include a means for injecting chemicals or DI water in lieu of the ozone-water solution and thereby provide a means to selectively dispense chemicals or DI water onto the surface of the materials to be oxidized during a given phase materials processing cycle

### Wafer Processing Advantages

higher removal rates: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like which can produce much higher removal rates than can be achieved by current methods

lower process temperature and lower corrosion potential: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like which can produce much higher removal rates at lower temperatures than can be achieved by current methods and thereby reduce the potential for metal corrosion

practical throughputs in both single wafer and batch processing systems: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like which can achieve practical throughputs in both single wafer processing systems and batch wafer processing systems

readily retrofitted at low marginal cost: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like which can be readily retrofitted at low marginal cost to many existing single wafer and batch wafer spin processing tools

readily integrated into a cluster tool: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like which can be readily integrated into a cluster tool comprising multiple processes in a single platform

readily integrated with a spin rinse and spin dry step: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like which can be readily integrated with a spin rinse and spin dry step provides the basis of a dry-in dry-out cleaning process

low cost processing chamber: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like which does not require that high concentration ozone gas be introduced into the process chamber thereby significantly reducing the cost of the wafer processing chamber.

increased user safety: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like which does not require that high concentration ozone gas be introduced into the process chamber thereby significantly increasing user safety

reduced corrosion potential: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like which does not require that high concentration ozone gas be introduced into the process chamber thereby eliminating high concentration ozone gas as a source of corrosion

nitrogen blanketed process chamber: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like in which an inert gas such a nitrogen can be introduced into the process chamber

increased user safety: provides a method for removing photoresist, post ash photoresist residue, post-etch residue, and other organic materials from semiconductor wafers, flat panel display substrates, and the like in which the dissolved ozone concentration quickly falls to

very low levels immediately downstream of the point of application thereby further increasing user safety

### Medical Instrument Sterilization Advantages

higher sterilization rates: provides a method for sterilization of medical instruments at moderate temperatures which can achieve much higher sterilization rates than can be achieved by current methods

residue free sterilant: provides a method for sterilization of medical instruments at moderate temperatures which utilizes a residue free sterilant and thereby eliminate the risk of carry-over of sterilant chemical residue to the patient and reduce the cost of chemical disposal

no water rinse required: provides a method for sterilization of medical instruments at moderate temperatures which utilizes a residue free sterilant and thereby eliminate the need for a separate water rinse step.

increased user safety and reduced chemical cost and reduced chemical disposal cost: provides a method for sterilization of medical instruments at moderate temperatures in which the active component is created for each cycle and then destroyed at end of the cycle thereby increasing user safety, decreasing chemical cost, and decreasing chemical disposal cost

lower consumables cost: provides a method for sterilization of medical instruments at moderate temperatures which has a lower consumables cost (\$.50 per cycle) than leading processes.

single-use sterilant always at full concentration: provides a method for sterilization of medical instruments at moderate temperatures which is a single pass design in which sterilant is sprayed onto instrument surfaces and then sent to the drain thereby eliminating the degradation in the sterilant concentration otherwise caused by the retention in the sterilant solution of serum and other organic residue washed from the processed instruments

low cost sterilization processing chamber: provides a method for sterilization of medical instruments at moderate temperatures which does not require that high concentration ozone gas be introduced into the process chamber thereby significantly reducing the cost of the wafer processing chamber.

increased user safety: provides a method for sterilization of medical instruments at moderate temperatures which does not require that high concentration ozone gas be introduced into the process chamber thereby significantly increasing user safety

reduced potential for instrument materials degradation: provides a method for sterilization of medical instruments at moderate temperatures which does not require that high concentration

ozone gas be introduced into the process chamber thereby eliminating high concentration ozone gas as a source of corrosion of metals or degradation of elastomers or plastics.

reduced potential for instrument materials degradation provides a method for sterilization of medical instruments at moderate temperatures in which an inert gas such a nitrogen can be introduced into the process chamber thereby eliminating high concentration ozone gas as a source of corrosion of metals or degradation of elastomers or plastics.

increased user safety: provides a method for sterilization of medical instruments at moderate temperatures in which the dissolved ozone concentration quickly falls to very low levels immediately downstream of the point of application thereby further increasing user safety

applicable to complex shaped instruments with internal passages with large L/D: provides a method for sterilization of medical instruments at moderate temperatures which can be used with complex shaped items, or items containing internal surfaces, such as rigid and flexible endoscopes with internal passages with a large length to diameter ratio L/D.

#### BRIEF DESCRIPTION OF DRAWINGS

The various features of the present invention and its preferred embodiments may be better understood by referring to the following discussion and the accompanying drawings in which like reference numerals refer to like elements in the several figures.

Fig. 1 illustrates a functional block diagram of a method of processing materials: A cold ozone-water solution at temperature  $T_1$  is heated to temperature  $T_2 > T_1$  using a liquid to liquid heat exchanger just upstream of the point of application of the ozone-water solution to the material to be processed.

Fig. 2 illustrates a functional block diagram of a method of processing materials: A cold ozone-water solution at temperature  $T_1$  is heated to temperature  $T_2 > T_1$  using a point-of-use heater just upstream of the point of application of the ozone-water solution to the material to be processed.

Fig. 3 illustrates a functional block diagram of a method of processing materials in which additional chemicals are injected into the ozone-water solution just upstream of the point of application.

Fig. 4 illustrates a functional block diagram of a method of processing materials with multiple chemical injection supplies.

Fig. 5 illustrates a block diagram of a method of processing semiconductor wafers with a single-wafer spin processor.

Fig. 6 illustrates a functional block diagram of a method of processing materials with upper and lower rotating spray arms (dishwasher geometry).

Fig. 7 illustrates a functional block diagram of a method of processing materials with an immersion design.

Fig. 8 illustrates a block diagram of a method of processing semiconductor wafers with multiple single-wafer spin processors - multiple point-of-use heater.

Fig. 9 illustrates a block diagram of a method of processing semiconductor wafers with multiple single-wafer spin processors - single point-of-use heater.

Fig. 10 illustrates a block diagram of a method of processing a batch (one or more cassettes) of semiconductor wafers with an on-axis batch spin processor.

Fig. 11 illustrates a block diagram of a method of processing a batch (one or more cassettes) of semiconductor wafers with an off-axis batch spin processor.

Fig. 12 illustrates a functional block diagram of a method of processing materials: A cold ozone-water solution at temperature  $T_1$  is applied to a material to be processed while the material to be processed and the ozone-water solution is heated to temperature  $T_2 > T_1$  at the point of application using radiant heating - heated from top surface.

Fig. 13 illustrates a functional block diagram of a method of processing materials: A cold ozone-water solution at temperature  $T_1$  is applied to a material to be processed while the material to be processed and the ozone-water solution is heated to temperature  $T_2 > T_1$  at the point of application using radiant heating - heated from bottom surface.

Fig. 14 illustrates a functional block diagram of a method of processing materials: A cold ozone-water solution at temperature  $T_1$  is applied to a material to be processed while the material to be processed and the ozone-water solution is heated to temperature  $T_2 > T_1$  at the point of application using a convection heating - heated from bottom surface.

Fig. 15 illustrates a functional block diagram of a method of processing materials: A cold ozone-water solution at temperature  $T_1$  is applied to a material to be processed while the material to be processed and the ozone-water solution is heated to temperature  $T_2 > T_1$  at the point of application using conduction heating - heated from bottom surface.

Fig. 16 illustrates a functional block diagram of a method of processing materials: A cold ozone-water mist solution at temperature  $T_1$  mixed with ozone gas, is applied to a material to be processed while the material to be processed and the ozone-water mist solution is heated to temperature  $T_2 > T_1$  at the point of application using radiant heating - heated from top surface.

Fig. 17 illustrates a block diagram of a method of dissolving ozone gas into chilled water using a venturi injector and downstream bubble column to form an ozone-water solution.

Fig. 18 illustrates a block diagram of a method of dissolving ozone gas into chilled water using a packed column to form an ozone-water solution.

Fig. 19 illustrates a block diagram of a method of dissolving ozone gas into water chilled water mist to form an ozone-water mist solution mixed with ozone gas.

## DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

### FACTORS DETERMINING OXIDATION RATE OR REMOVAL RATE - A MODEL

The inventors have developed a model to help better understand the factors determining oxidation and removal rate of an organic material such as photoresist from a semiconductor wafer using an ozone-solvent solution at concentration  $C$  and temperature  $T$ . The rate of oxidation and removal of an organic layer from a substrate can be defined in terms of an etch rate. We can write an expression for the etch rate  $E$  (cm resist/sec) as

$E = C * (X/\rho) * (M * S) / (M + S)$ . The parameter  $C$  (g ozone/cm<sup>3</sup>) is the dissolved ozone concentration in the water far from the surface of the organic layer on a semiconductor wafer for example (the bulk concentration). The parameter  $X$  (g resist/g Ozone) is the mass of resist removed per mass of ozone consumed at the surface. If we assume that the resist is composed of chains of CH<sub>2</sub> units which must be fully oxidized to be removed, then 3 moles of ozone are required to oxidize each mole of CH<sub>2</sub>. This corresponds to 10.3 grams of ozone to fully oxidize each gram of resist. Christensen (1998) was the first to observe that etch rates for I-line photoresist are a factor of 20 higher than that predicted by the assumption of full oxidation. They concluded that the resist need only be cut into short fragments about 20 CH<sub>2</sub> units long before the fragments become hydrophilic and float off the wafer into the stream of flowing water. The net result is that the parameter  $X$  is not (1/10.3) but about (20/10.3). The parameter  $\rho$  (g resist/cm<sup>3</sup>) is the density of the resist.

Terminology: Throughout this discussion the terms ozonated water, ozone-water solution, and ozone-gas-water solution are used interchangeably. In addition, the terms etch, etch clean, clean, process, oxidize are used interchangeably. DI water is De-ionized water

**Dissolved Ozone Concentration C:** When ozone is dissolved in a solvent, the maximum dissolved ozone concentration C that can be achieved after a sufficiently long transfer time, the saturation concentration, is predicted by Henry's law. According to Henry's law, the maximum solubility is proportional to the partial pressure of the ozone gas at a given temperature. Higher gas phase concentrations, high pressures, and lower solvent temperatures yield higher maximum equilibrium dissolved ozone concentrations. We have calculated the approximate equilibrium saturation concentration in mg/L (equivalent to parts per million by weight) for a gas phase concentration of 240 mg/L (15.9 weight percent), pressures of 1, 2, and 4 bar, and water (solvent) temperatures of 5 to 95 degree C in 5 deg. C increments. See Table 1.

TABLE 1. Solubility of ozone gas in water: The dissolved ozone concentration in mg/L as a function of the water temperature and gas pressure for a gas phase ozone concentration of 240 g/Nm<sup>3</sup>=mg/liter (15.9 weight percent) in oxygen for a range of water temperatures.

	p=1 bar (14.5 psia)	p=2 bar (29 psia)	p=4 bar (58 psia)
5 deg. C	109	218	436
10 deg. C	85	170	340
15 deg. C	66	132	264
20 deg. C	52	104	208
25 deg. C	40	80	160
30 deg. C	31	62	124
40 deg. C	24	48	96
45 deg. C	19	38	76
50 deg. C	15	30	60
55 deg. C	11	22	44
60 deg. C	9	18	36
65 deg. C	7	14	28
70 deg. C	5	10	20
75 deg. C	4	8	16
80 deg. C	3	6	12
85 deg. C	2.5	5	10
90 deg. C	1.2	2.4	4.8
95 deg. C	.9	1.8	3.6

Mass Transport Rate Coefficient M: The parameter  $M$  (cm/sec) is the liquid phase mass transport rate coefficient. The ozone is transported to the wafer surface by diffusion. The mass transport rate  $M$  (cm/sec) =  $D/\delta$ , where  $D$  (cm<sup>2</sup>/sec) is the diffusion constant of the ozone diffusing in the liquid and  $\delta$  (cm) is the thickness of the stagnant layer. The diffusion constant  $D$  for ozone in water is  $1.7E-5$  cm<sup>2</sup>/sec at 20 deg. C. Accordingly the mass transport rate is increased when the diffusion constant is increased and/or the diffusion distance  $\delta$  is decreased.

Surface Reaction Rate Constant S: The parameter  $S$  (cm/sec) is the temperature dependent surface reaction rate constant. The surface reaction rate  $S$  (cm/sec) is an exponential function of the absolute temperature  $T$  (deg. K) and the activation energy  $E_a$  of the oxidation process. In particular,  $S = S_0 \exp(-E_a/KT)$  where  $K$  is Boltzman's constant and  $S_0$  is the surface reaction rate proportionality constant. The difference in etch rates of different materials at a given temperature is attributed to the difference in the magnitude of the surface reaction rate constant for the two materials.

Etching Wafers at High Temperature: An increase in temperature will increase  $S$  and the magnitude of the term  $(M*S)/(M+S)$ . If the dissolved concentration remained approximately constant with an increase in temperature, then we can see that the etch rate would increase with increased temperature. However, as we have seen, the dissolved ozone concentration falls with increases in water temperature. If the temperature is such that  $S \gg M$ , the etch rate becomes mass transport limited and  $E = C(X/\rho)*M$ . If  $M$  is larger, then the temperature at which the etch rate becomes limited by the mass transport rate  $M$  will be higher. If the temperature is higher, then the mass transport rate at which the etch rate will become mass transport limited will be higher.

Etching Wafers at Low Temperature: A decrease in temperature will decrease  $S$  and the magnitude of the term  $(M*S)/(M+S)$ . If the dissolved concentration remained approximately constant with decreases in temperature, then we can see that the etch rate would decrease with a decrease in temperature. However, as we have seen, the dissolved ozone concentration rises with decreases in water temperature. If the temperature is decreased until  $S \ll M$ , the etch rate becomes surface reaction rate limited and  $E = C(X/\rho)*S$ .

Measured Temperature Dependence of S: We made a preliminary measurement of the temperature dependence of the surface reaction rate  $S$  in our laboratory. We made the



measurements of the etch rate of IBM Apex positive DUV resist under conditions where  $M \gg S$  and  $E = SCX$  by etching a spinning wafer with a small diameter nozzle to produce high velocities and small values of  $\delta$ . We measured the etch rate at 7 deg. C, 12 deg. C, and 17 deg. C. Our measurements showed that the surface reaction rate coefficient  $S$  (cm/sec) increases by a factor of two for every 5 degree C increase in temperature corresponding to an activation energy of about 1 eV. We found the same activation energy but larger coefficient  $S_0$  for Shipley 1805 positive I-line resist. This is consistent with I-line positive resist etching faster than DUV positive photoresist.

Measured Etch Rate at Different Temperatures: Our initial measured results show that the etch rate increases when the temperature is increased from 10 deg. C to 19 deg. C because the normalized etch rate  $E/C$  increases by a larger factor than the dissolved ozone concentration decreases. However, when the temperature is increased from 19 deg. C to 28 deg. C, the dissolved ozone concentration decreases by a larger factor than the normalized etch rate decreases. This trend continues at all higher temperatures with the result that under equilibrium conditions the highest etch rate is achieved at about 20 deg. C.

An Approach to Achieving Very High Etch Rates: This model can provide us valuable insight into the problem. It shows that the normalized etch rate be increased by increasing the temperature and that the etch rate could be increased by increasing the temperature above 20 degree C if we could find a method to provide a higher dissolved concentration at the elevated temperature. The present preferred embodiments utilize just such a method.

The general principal is to achieve the highest dissolved ozone concentration at a given surface reaction temperature. This can be done in a number of ways including the following:

- a) heat the cold ozone-solvent solution with an in-line heater located just upstream of the point at which the ozone-solvent solution is dispensed onto the substrate. The heated ozone-solvent solution will then heat the surface of the substrate and increase the surface reaction rate. The ozone-solvent solution will retain most of the ozone dissolved at the lower temperature if the solution is not heated until the last moment.
- b) heat the cold ozone-solvent solution at the point of application with a point of application heater as the solution passes over the substrate surface by for example using a radiant heater with the wavelength band chosen to be absorbed by the ozone-solvent solution. The heated ozone-solvent solution will then heat the surface of the substrate and increase the surface reaction rate.

- c) heat the substrate with a point of application heater and dispense the cold ozone-solvent solution onto the surface of the heated substrate. Provide sufficient heat input to the substrate to overcome cooling effect of the cold solvent. In practice the substrate can be heated from the backside or from the front side. If the substrate is heated from the backside, the entire volume of the substrate may be heated so that the front surface, the surface to be etched, can be heated. If the substrate is heated from the front side, the entire volume of the substrate may be heated or only the front surface may be heated. The surface reaction rate at the front surface is a function of the temperature of the front surface.
- d) heat the cold ozone-solvent solution and heat the substrate by for example using a radiant heater with the wavelength band chosen to be absorbed partially by the ozone-solvent solution and partially by the substrate.

The substrate surface can be heated by conduction, convection, or radiation. The surface can be heated by conduction using a heated surface such as a hot plate. The surface can be heated by convection using a hot gas or hot liquid to the front or rear surface. The substrate can be heated by radiation using a heat lamp or laser or other source of radiation. The radiation wavelength band can be chosen so that the radiation passes through the ozone-water solution with little energy deposition in the water and the majority of the energy absorbed in the surface. In fact the radiation can be chosen to be most strongly absorbed in the layer to be removed (photoresist for example).

The description of the preferred embodiments for oxidizing materials at high speed is divided into three groups

1st GROUP: Heat Applied Just Upstream of the Point of Application (Fig. 1 through Fig 11)

2nd GROUP: Heat Applied at the Point of Application (Fig. 12 through Fig 16)

3rd GROUP: High Performance Ozonated Water Supply for Use in Conjunction with the Preferred Embodiments (Fig. 17 through Fig. 19)

1st GROUP: Heat Applied Just Upstream of the Point of Application (Fig. 1 through Fig 11)

A first preferred method for oxidizing materials at high speed using a solution of ozone gas dissolved in solvent comprising the steps of dissolving relatively high concentration ozone gas in solvent at a relatively low predetermined temperature  $T_1$  to form an ozone-solvent solution with a relatively high dissolved ozone concentration, and heating the ozone-solvent solution with a point-of-use heater to quickly increase the solution temperature to a predetermined higher temperature  $T_2 > T_1$ , and applying the heated ozone-solvent solution to said material(s). The first method for oxidizing materials at high speed may additionally

includes an injector/mixer for injecting and mixing additional chemicals into the ozone-solvent solution just upstream of the point of application of the ozone-solvent solution to the materials to be oxidized. A number of preferred embodiments are illustrated in Fig. 1 through Fig. 11.

#### Materials Processing Method W/ Ozone-Solvent Solution Heated W/ Heat Exchanger

Description - Fig. 1 With reference to Fig 1, an ozonated water supply 22 is connected through a length of tubing to the common input of a three-way valve 24. The one outlet of three-way valve 24 is connected through a length of tubing to the cold process fluid inlet of a heat exchanger 28. The other outlet of three-way valve 24 is connected through a length of tubing to the facility drain-reclaim inlet 26 for the ozone-water solution. The heated working fluid outlet of a recirculating heating unit 30 is connected through a length of tubing to the heated working fluid inlet of heat exchanger 28. The heated working fluid outlet of heat exchanger 28 is connected through a length of tubing to the working fluid return of recirculating heating unit 30. The heated process fluid outlet of heat exchanger 28 is connected through a short length of tubing to a three-way valve 32. The one outlet of three-way valve 32 is connected through a short length of tubing to a dispense nozzle 36. The dispense nozzle 36 is spaced a relatively short distance from the surface 38 of the material to be oxidized, cleaned, or processed. The other outlet of three-way valve 32 is connected through a length of tubing to the facility drain-reclaim inlet 34 for the heated ozone-water solution.

#### Operation - Fig. 1

With reference to Fig 1, an ozone-gas-water solution of predetermined dissolved ozone concentration is produced by the ozonated water supply 22 by dissolving ozone gas at predetermined concentration and pressure P1 into water at a predetermined temperature T1. The ozone gas is dissolved into water using a venturi injector and downstream bubble column, a packed column, a gas permeable membrane contactor, a bubble diffuser, a turbine mixer, a spray contactor, or other means known to those skilled in the art. The ozone gas may be dissolved at atmospheric pressure (~ 1 bar) or at elevated pressures above atmospheric pressure. The maximum dissolved concentration that can be produced for a given gas phase concentration, gas pressure, and water temperature is predicted by Henry's law as discussed in the introduction. (See Table 1).

Some ozone generators such as the Astex AX8100 and AX 8200 require high purity oxygen gas mixed with approximately 0.5 % nitrogen gas by volume to produce relatively high concentration ozone gas (200 to 250 grams/Nm<sup>3</sup>). Other generators such as the Astex

Model AX8401, Astex Model AX8402, Semozon Model 030.2, and Semozon Model 090.2 require a source of high purity oxygen with only about 50 ppm of nitrogen by volume to produce relatively high concentration ozone gas. A generator requiring oxygen mixed with approximately 0.5% nitrogen produces ozone ( $O_3$ ) and a relatively large amount of  $NO_2$ . When this gas stream is dissolved in water in the ozone-water contactor, the  $NO_2$  combines with water ( $H_2O$ ) to form nitric acid ( $HNO_3$ ). If the water is unbuffered, then nitric acid can cause the pH of the ozone-water solution to gradually fall. In the preferred electronic device cleaning/processing embodiment, the use of an ozone generator requiring oxygen mixed with approximately 50 ppm nitrogen is preferred because this type of generator produces ozone ( $O_3$ ) and a very small amount of  $NO_2$ . Accordingly, when this gas stream is dissolved in water in the ozone-water contactor, only a small amount of nitric acid is formed, and there is a minimal resulting pH change.

The ozonated water supply 22 may be designed to supply an ozone-gas-water solution at a predetermined concentration and flow rate on a continuous basis or on an intermittent basis. In general, the mass flow rate out of the ozone-water solution (the product of the liquid flow rate and the dissolved ozone concentration) is less than the mass flow rate of the ozone gas entering the contactor (the product of the ozone gas flow rate and the ozone gas concentration). Accordingly, some of the ozone gas will exit from the contactor vent line as a waste ozone gas stream.

An ozonated water supply may be designed to flow the water through the gas-water contactor in a single pass and deliver an ozone-gas-water solution at a predetermined value less than the saturation concentration. Alternatively, the ozonated water supply may be designed to flow the water through the gas-water contactor in multiple passes, and thereby provide a longer time for mass transfer to occur between the gas and the liquid, and deliver an ozone-gas-water solution at a predetermined value up to the saturation concentration.

It is convenient to use a three way valve 24 to direct the solution to the process for the required process cycle time, and then direct the ozone-water solution to facility waste/reclaim 26 at the end of the cycle. This is preferred to shutting off the flow for ozonated water supplies based upon a single pass design since this type of ozonated water supply 22 requires water flow the operate. This insures that the supply runs in steady state and can provide a stable supply of ozonated water at a predetermined concentration with no transients associated with stopping and starting flow through the supply contactor. The supply can employ a controller to set the flow to a very low level during the period that the valve is set to the waste-reclaim position to conserve water and ozone. If an ozonated water supply of the batch

recirculating type is employed, then the three-way valve may be used to direct the flow of water back to the contactor.

If the flow through heat exchanger 28 is stopped at the end of a process cycle, then the dissolved ozone concentration in the water remaining in the exchanger will decay toward a low value and the temperature will rise to the hot working fluid temperature. It is convenient to use a three way valve 32 just upstream of the dispense nozzle 36 to direct this volume of water to facility waste/reclaim 34 immediately prior to the next process dispense cycle. This insures that the heated ozone-gas-water solution that is subsequently dispensed onto the material 38 is at the predetermined temperature and dissolved ozone concentration. At the start of an etch cycle (process cycle) valve 26 is set to direct the process flow through the point-of-use heater (heat exchanger 28) and valve 32 is set to direct the process flow to facility waste/reclaim 34 for a time greater than or equal to the residence time of the volume between three-way valve 24 and three-way valve 32. In the case of a flow rate of 50 ml/sec (3.0 L/min) and a residence volume of 100 ml, the flow is sent to facility waste/reclaim 34 for a least 2 seconds to insure that the ozone-water solution is purged from the residence volume before valve 32 is set to direct the process flow to the dispense nozzle 36. The valve 32 is placed just upstream of the dispense nozzle 36 to minimize the volume that is not purged when valve 32 is set to direct the process flow to facility waste/reclaim 34. The waste-reclaim 26 and waste-reclaim 34 may be directed to different facility waste-reclaim locations since the process stream sent to 26 is the full concentration ozone-gas-water solution supplied by the ozonated water supply 22 and the process stream sent to 34 is the heated ozone-gas-water solution in which the concentration may have fallen significantly during the time when the flow through exchanger 28 has been set to zero between the end of one process cycle and the start of the next process cycle.

The ozonated water supply 22 delivers an ozone-gas-water solution with sufficient pressure to achieve a predetermined flow rate through the dispense nozzle 36. The delivery pressure must be sufficient pressure to produce the flow through the pressure drop across the heat exchanger 28, connecting tubing, and dispense nozzle 36.

Back Pressure Regulators for an Ozonated Water Supply Using a Pressurized Contactor. If the ozonated water supply 22 is designed to dissolve ozone gas into water at a gas pressure P2 above 1 bar, then a back pressure regulator is typically placed at the ozone off-gas outlet of the ozone-gas-water contactor and ozonated water outlet of the ozone-gas-water contactor to maintain the specified gas pressure P2 inside the contactor which is higher than pressure P1 downstream of the back pressure regulators. Once the ozone-water solution

passes through the back pressure regulator to a lower pressure the ozone gas will begin to leave the solution. If the transit time of the ozone-gas-water solution from the outlet of the ozone-gas-water contactor in the ozonated water supply 22 to the inlet of the heat exchanger 28 is relatively long, then the ozone gas will have a longer time available to leave the solution in transit from the back pressure regulator to the exchanger inlet. The dissolved ozone concentration at the exchanger inlet will be less than the dissolved ozone concentration at the outlet of the ozonated water supply 22 just downstream of back pressure regulator because the ozone concentration will fall toward the equilibrium concentration at the pressure downstream of the regulator. This fall in concentration can be alleviated by moving the back pressure regulator to a point just upstream of heat exchanger 28. If the back pressure regulator is moved further downstream to a point just upstream of the dispense nozzle 36, then the decay in concentration caused by the  $P_2 > P_1$  can be reduced even further since the solution is maintained at the pressure  $P_1$  until last possible moment when the solution is dispensed onto the surface of the material 38 at a pressure  $P_2$ .

Heat Exchanger : A Point-of-Use Heater for Quickly Heating the Ozone-Gas-Water Solution. The ozone-water solution flows through three-way valve 24 to the process fluid inlet of the heat exchanger 28. A heated working fluid such as water, supplied by a recirculating heating unit 30, flows through the hot working fluid side of the heat exchanger. The recirculating heating unit 30 is sized to provide the power required to increase the ozone water solution temperature at a specified ozone water solution flow rate. The temperature hot working fluid (typically water) that flows from the recirculating heating unit 30, through the exchanger 28, and back to the recirculating heating unit 30 for reheating is controlled by the temperature controller of the recirculating heating unit. The temperature of the ozone water solution exiting from the exchanger can be changed by changing the temperature setpoint on the recirculating heating unit. In passing through the heat exchanger, the ozone-water solution is quickly heated to a higher temperature and then immediately applied to the material to be oxidized.

The volume of the fluid passages through which the heated, supersaturated, ozone-water solution must pass, starting at the inlet of the heat exchanger 28 and ending at the dispense nozzle 36 must be kept relatively small. A typical value for this volume is 100 to 300 ml for a system designed for a dispense flow rate of approximately 3.0 l/min and a dispense temperature of approximately 50 degree C. This corresponds to a residence time of 2 seconds to 6 seconds. In addition, the volume of the fluid stream starting at the dispense nozzle 36 and ending at the surface of the material to be oxidized 38 must be kept small. A

typical value for this volume is 5 to 10 ml for a system designed for a dispense flow rate of approximately 3.0 l/min and a dispense temperature of approximately 50 degree C. This corresponds to a residence time of .1 seconds to .2 seconds. A minimum total residence time, the time delay between heating the solution and applying the solution to the surface(s) to be cleaned or oxidized, minimizes the time available for the dissolved ozone concentration to fall once the solution temperature is increased.

The ozone gas will come out of solution at a higher rate and the dissolved ozone concentration will fall at a higher rate if the ozone-water solution is heated to a higher temperature. The optimum temperature for maximizing the etch rate is determined by the mass transport coefficient  $M$  as discussed earlier. Once  $S \gg M$  at sufficiently high temperature, further increases in temperature do not increase etch rate further. One design approach is then to design the point-of-use heater to have a sufficiently small residence time at that etch rate maximizing temperature that the dissolved ozone concentration decays a relatively small amount.

The ozone gas will come out of solution at a higher rate and the dissolved ozone concentration will fall at a higher rate if the inside surface of the point-of-use heater (the process side of the heat exchanger in this embodiment) has scratches, since the ozone will leave solution most readily at the location of the scratches. Accordingly, a relatively smooth inner surface is desired. The inside surface of the 316 stainless steel tube-in-tube heat exchanger model 413 made by Exergy Inc. has a surface roughness of  $\sim 20$  Ra. Electropolished versions are available with a surface roughness of  $\sim 5$  Ra.

The ozone gas will come out of solution at a higher rate and the dissolved ozone concentration will fall at a higher rate if the surface area of the free stream(s) of fluid passing from the dispense nozzle(s) to the surface 36 is larger. If the flow of the solution from the dispense nozzle to the surface of the material to be oxidized 38 is carried by many small diameter streams, then the rate of loss of ozone gas from solution will be higher when compared to the case of the solution being carried by a single solid stream because the surface area of the many small diameter streams is larger.

In applications such as wafer processing the use of stainless steel wetted components is not acceptable because these materials introduce metal contamination into the process chemistry. In these applications the preferred materials for all wetted parts are metal free materials such as Teflon PFA or Teflon PTFE or quartz. Accordingly, the point-of-use heater should employ materials such as Teflon PFA or Teflon PTFE for all wetted surfaces in lieu of stainless steel for these applications.

In the prior art, the ozone is dissolved in water at a given temperature  $T_1$  and the ozone-water solution is applied to the material to be oxidized at a temperature  $T_2=T_1$ . Accordingly, if one lowers the temperature  $T_1$  to increase the dissolved ozone concentration, then one decreases the surface reaction rate  $S$ . Alternatively, if one increases the temperature  $T_2$  to increase the surface reaction rate, then one decreases the dissolved ozone concentration.

The present invention overcomes this limitation by tricking mother nature into providing a higher dissolved concentration at the elevated temperature that could be achieved under equilibrium conditions.

A first type of method for treating materials at high speed using a solution of ozone gas dissolved in a solvent comprises the steps of dissolving relatively high concentration ozone gas in water at a relatively low predetermined temperature  $T_1$  to form an ozone-water solution with a relatively high dissolved ozone concentration, and heating the ozone-water solution with a point-of-use heater to quickly increase the solution temperature to a predetermined higher temperature  $T_2>T_1$ , where preferably  $T_2-T_1>5$  degree C, and applying the heated ozone-water solution to said material(s).

A second type of method for oxidizing materials at high speed using a solution of ozone gas dissolved in solvent comprises the steps of dissolving (relatively high concentration) ozone gas in water at a relatively low predetermined temperature  $T_1$  to form an ozone-water solution (with a relatively high dissolved ozone concentration), applying the cold ozone-water solution to said materials while heating said materials and said ozone-water solution at the point of application to quickly increase said material temperature and said solution temperature to a predetermined higher temperature  $T_2>T_1$ , where preferably  $T_2-T_1>5$  degree C.

The heated ozone-water solution will have a much higher dissolved ozone concentration at said higher temperature than could be achieved if the ozone gas was initially dissolved in water at said higher temperature. The parameter space for the preferred embodiments and prior art is shown in Table 2 below. In the prior art, the ozone is dissolved and applied at the same temperature.



Table 2: Parameter Space of the Preferred Embodiments and Prior Art (the ozone-water solution is prepared by dissolving ozone gas in water at a temperature  $T_1$ ; the ozone-water solution is applied to the material to be oxidized at a temperature  $T_2$ )

$T_2 > T_1$	Preferred Embodiments
$T_2 = T_1$	PRIOR ART
$T_2 < T_1$	generally lower performance than $T_2 = T_1$ (Note 1)

Note 1: In the case with  $T_2 < T_1$  the dissolved concentration is nominally the same as when  $T_2 = T_1$  and the surface reaction rate  $S$  is lower because the surface temperature is lower.

The point-of-use heater is designed to have a small residence volume so that the residence time between the inlet of the heater and the point of application is small and there is insufficient time for supersaturated solution to return to equilibrium before reaching the surface of the material to be oxidized. The time required for the solution to return to equilibrium is dependent upon the temperature to which the solution is heated. Our preliminary measurements indicate that at a temperature of about 50 degree C, a residence time of 2 seconds will allow the dissolved concentration to only fall by about 10 to 20 percent. At higher temperatures, the required residence time is smaller. The residence time is proportional to the volume and inversely proportional to the dispense flow rate though that volume.

We prepared an ozone-water solution by dissolving ozone gas, at concentration of 240 g/Nm<sup>3</sup>, a flow rate of 0.48 L/min, and a pressure of 1 bar, into water at a temperature of about 8 degree C using a Mazzei Model 287 venturi injector and bubble column contactor operated in the recirculating mode. We waited about 30 minutes and allowed the dissolved concentration reach the saturation concentration at about 70 . We drew the ozone-water solution from the unpressurized contactor with a high pressure gear pump capable of delivering a flow rate of 2.7 L/minute at 80 psi. We passed the solution through an Exergy tube in tube heat exchanger model 413, through a UV absorption type dissolved ozone monitor and thermocouple probe, and then to a waste collection carboy. We measured the dissolved concentration upstream and downstream of the heater as a function of the temperature of the ozone-water solution downstream of the heater for several different temperatures. We used this data to estimate the decay time constant as function of temperature by assuming that the decay time was an exponential function of the temperature. We ran a similar test using coil in heated water bath heater. We flowed the water through a

foot long coil of stainless steel, .375 inch OD, .305 inch ID tubing, immersed in a heated water bath. Since the water bath did not have sufficient power to maintain a constant bath temperature, the dispense temperature of the ozone water solution decreased about 5 degree C during the test. Accordingly, we used average temperatures in analyzing the results. The residence volume of the coil in bath heater was about 270 ml and the residence volume of the tube in tube heat exchanger was about 90ml. (see the table 3 footnotes) The results for both tests were consistent with the model presented below. The results for one test series are presented in Table 3 below.

Table 3. Decay time constant as a function of temperature: Measured decay time constant as function of temperature and calculated decay time constant as function of temperature assuming that the decay time is an exponential function of the temperature

Tau=2E-10*Exp(8.26(1000/T))									
		measured		calculated					
		decay		decay		transit		transit	
		const.		const.		time		time	
Temp	Temp	Tau		Tau		t		decay t	
deg.	deg.	1000/T		secs		secs		exp(-	
C	K	(K)		secs		exp(-		t/Tau	
						t/Tau)		)	
						note 1		note 2	
								note 3	
20	293	3.41		292.3	2	99%	6	98%	22
25	298	3.36		186.6	2	99%	6	97%	22
30	303	3.30		120.9	2	98%	6	95%	22
35	308	3.25		79.4	2	98%	6	93%	22
40	313	3.19		52.9	2	96%	6	89%	22
45	318	3.14		35.7	2	95%	6	85%	22
50	323	3.10		24.3	2	92%	6	78%	22
55	328	3.05		16.8	2	89%	6	70%	22
60	333	3.00	11.00	11.7	2	84%	6	60%	22
65	338	2.96	8.4	8.3	2	79%	6	49%	22
70	343	2.92		5.9	2	71%	6	36%	22

75	348	2.87	4.05	4.3	2	63%	6	24%	22	1%
80	353	2.83		3.1	2	52%	6	14%	22	0%
85	358	2.79		2.3	2	41%	6	7%	22	0%
90	363	2.75		1.7	2	30%	6	3%	22	0%
95	368	2.72		1.3	2	20%	6	1%	22	0%

Note 1: Exergy Model 413 stainless steel tube-in-tube heat exchanger; total residence volume from the heater inlet to the heater outlet approximately 90 ml; dispense flow rate 45 ml/sec; residence time approximately 2 secs

Note 2: 3/8 inch OD, 20 foot long, coil in water bath heater; total residence volume from the heater inlet to the heater outlet approximately 270 ml; dispense flow rate approx. 45 ml/sec; residence time approximately 6 secs

Note 3: relatively large volume heater or heat exchanger; total residence volume from the heater inlet to the heater outlet chosen to be 1000 ml; dispense flow rate approx. 45 ml/sec; residence time approximately 22 secs

From this data we can see that higher temperatures cause the concentration to fall more quickly. If one is to minimize the drop in concentration upon heating, then the residence time must be decreased if the temperature is increased. For example, if we would like the dissolved ozone concentration at the outlet of the point-of-use heater to be no less than 80 percent of the dissolved ozone concentration at the inlet of the point-of-use heater, then the transit time must be less than or equal to the values estimated in Table 4 below.

Table 4. Maximum estimated permissible ozone-water solution heating time (heater transit time): Example calculated for the dissolved ozone concentration at the heater outlet to be no less than 80 percent of the dissolved ozone concentration at the heater inlet. Estimated from decay data measured with an inlet dissolved ozone concentration of about 100 mg/liter, an initial upstream ozone-water solution temperature of about 8 deg. C and the a final specified downstream ozone-water solution temperature ranging from 20 deg. C to 95 deg. C

Table 4.

Example Decay Factor = 80% (Downstream Conc./Upstream Conc.)			Estimated	Calculated
			decay	maximum allowable
Ave solution temp. at heater outlet			const.	heater transit time
Temp	Temp.		Tau	$t = -\text{Tau} * \ln(\text{decay factor})$
deg. C	deg. K	1000/T (K)	secs	secs
20	293	3.41	292.3	65.22
25	298	3.36	186.6	41.63
30	303	3.30	120.9	26.97
35	308	3.25	79.4	17.72
40	313	3.19	52.9	11.80
45	318	3.14	35.7	7.96
50	323	3.10	24.3	5.43
55	328	3.05	16.8	3.75
60	333	3.00	11.7	2.62
65	338	2.96	8.3	1.85
70	343	2.92	5.9	1.32
75	348	2.87	4.3	0.95
80	353	2.83	3.1	0.69
85	358	2.79	2.3	0.51
90	363	2.75	1.7	0.37
95	368	2.72	1.3	0.28

We know from our model that an increased temperature will provide a significant increase in etch rate until the etch rate becomes mass transport limited. Our measurements show that with a wafer spinning at about 4,000 rpm and a dispense flow rate of 2.7 L/min, that

the etch rate becomes mass transport limited at the wafer edge at about 50 degree C. Other methods of applying the solution to the material to be oxidized may provide a higher or lower mass transport coefficient and a correspondingly higher or lower optimum temperature.

**Heating Power Requirement.** The power input in watts required to increase the temperature of a stream of water can be calculated given the water flow rate, the desired temperature rise, and the heat capacity of the water. We have shown the results of that calculation in Table 5 below. If the flow rate of the ozone-water solution which passes through the point-of-use heater is 2.7 L/min, and the water enters at a temperature of 5 degree C, and the desired exit temperature is 55 deg. C, then the heater must transfer energy to the ozone-water stream at a power level of 9.3 kW.

Table 5. Point-of-use Heater Power Requirement

Dispense Flow Rate (L/min)	Inlet Water Temp. (deg. C)	Desired Outlet Water Temp. (deg. C)	Required Temp. Increase (deg. C)	Power Req'd (kW)
2.7	5	45	40	7.45
2.7	5	55	50	9.3
2.7	5	65	60	11.2
2.7	5	75	70	13.0
2.7	5	85	80	14.9
2.7	5	95	90	16.8
3.3	5	45	40	9.1
3.3	5	55	50	11.4
3.3	5	65	60	13.7
3.3	5	75	70	15.9
3.3	5	85	80	18.2
3.3	5	95	90	20.5

The power required to increase the ozone gas-water solution temperature can be transferred from a heated working fluid in a heat exchanger. The energy transferred by a particular heat exchanger is determined by the temperature and flow rate of hot working fluid entering the working fluid side of the exchanger, the temperature and flow rate of the ozone gas-water solution entering the process side of the exchanger, and the desired temperature of

the ozone-water solution exiting from the exchanger. Example flow rates and temperatures are shown in Table 6 below. We can see that for a given flow rate and temperature for the ozone-water solution entering the heat exchanger and a given flow rate for the heated working fluid circulated through the outer tube of the heat exchanger by the recirculating heating unit, that changing the temperature of the working fluid will change the temperature of the ozone-water solution exiting from the exchanger to the dispense nozzle. The embodiment can provide a constant dispense temperature T2 the other parameters mentioned are held constant.

Table 6: Calculated Performance of Exergy Model 413 Stainless Steel Tube-in-Tube Heat Exchanger Reported by the Manufacturer. The ozone-gas-water solution is passed through the 0.180 inch ID, .250 inch OD, 240 inch long, inner tube. The heated working fluid from the recirculating heating unit is passed through the annular space bounded by a concentric .430 inch ID outer tube and the inner tube. The internal volume of the inner tube is 90 ml.

	Example 1: 48 °C Dispense Temperature		Example 2: 56 °C Dispense Temperature	
	Inner Tube	Outer Tube	Inner Tube	Outer Tube
	ozone-water solution	heated working fluid	ozone-water solution	heated working fluid
Fluid Used for the Calculation	water	water	water	water
Temp In (°C)	7	60	7	70
Temp. Out (°C)	48	38	56	43
Vol. Flow (L/min)	2.7	5	2.7	5
Pressure Drop (psi)	24	6	24	6
Heat Transfer (watts)	7616	7616	9203	9203

Materials Processing Method W/ Ozone-Water Solution Heated W/ A Point-Of-Use Water Heater

### Description - Fig. 2

With reference to Fig 2, an ozonated water supply 22 is connected through a length of tubing to the common input of a three-way valve 24. The one outlet of three-way valve 24 is connected through a length of tubing to the cold process fluid inlet of point-of-use heater 29. The other outlet of three-way valve 24 is connected through a length of tubing to the facility drain-reclaim 26 for the ozone-water solution. The heated process fluid outlet of point-of-use heater 29 is connected through a short length of tubing to a three-way valve 32. The one outlet of three-way valve 32 is connected through a short length of tubing to a dispense nozzle 36. The dispense nozzle 36 is spaced a relatively short distance from the surface 38 of the material to be oxidized, cleaned, or processed. The other outlet of three-way valve 32 is connected through a length of tubing to the facility drain-reclaim 34 for the heated ozone-water solution.

### Operation - Fig. 2

**Point-of-Use Heater:** A Point-of-Use Heater for Quickly Heating the Ozone-Gas-Water Solution: The point-of-use heater 29 has the same requirements as the heat exchanger including those relating to internal volume, surface roughness, and materials of metal free materials of construction. Most commercially available heaters of the required power level and materials of construction such as those made for point-of-use heating of DI water for the semiconductor and pharmaceutical industry have an internal volume of at least 2000 ml. One embodiment for a point-of-use heater with small internal volume is constructed using radiant and convention heating of the ozone-water solution. The ozone-water solution is flowed through quartz tubing that surrounds the IR heating source. The volume of the tubing can be made small to minimize the residence time inside the heater. A temperature sensor, either at the outlet of the heater, or just upstream of the dispense nozzle, is connected to temperature controller. The temperature controller increases or decreases the amount of power delivered by the heater to achieve and maintain a specified dispense temperature. A direct heater with feedback control from the dispense temperature can typically adjust the temperature at the dispense point more quickly than can a heat exchanger embodiment.

**Heating Power Requirement:** The heating power requirement using a point-of-use heater 29 is the same as that using a heat exchanger 28.

### Materials Processing Method Additionally Including Point-Of-Use Chemical Injection - Single Chemical Supply-

#### Description- Fig. 3

It is often useful to have a means to mix additional chemicals with the ozone-gas-water solution prior to applying the solution to the material to be oxidized.

With reference to Fig 3, an ozonated water supply 22 is connected through a length of tubing to the common input of a three-way valve 24. The one outlet of three-way valve 24 is connected through a length of tubing to the cold process fluid inlet of a heat exchanger 28. The other outlet of three-way valve 24 is connected through a length of tubing to the facility drain-reclaim 26 for the ozone-water solution. The heated working fluid outlet of a recirculating heating unit 30 is connected through a length of tubing to the heated working fluid inlet of heat exchanger 28. The heated working fluid outlet of heat exchanger 28 is connected through a length of tubing to the working fluid return of recirculating heating unit 30. The heated process fluid outlet of heat exchanger 28 is connected through a short length of tubing to the inlet of chemical injector/mixer 40. The outlet of an injected chemical supply 42 is connected to the inlet of two-way valve 44. The outlet of valve 44 is connected to the chemical injection port of chemical injector/mixer 40. The outlet of chemical injector/mixer 40 is connected to the common inlet port of three-way valve 32. The one outlet of three-way valve 32 is connected through a short length of tubing to a dispense nozzle 36. The dispense nozzle 36 is spaced a relatively short distance from the surface 38 of the material to be oxidized, cleaned, or processed. The other outlet of three-way valve 32 is connected through a length of tubing to the facility drain-reclaim 34 for the heated ozone-water solution.

Operation - Fig. 3

Injected Chemical Supply System - Single Chemical Reservoir: The injected chemical supply provides a chemical at a predetermined delivery pressure and a predetermined flow rate into the injection port of the injector/mixer 40 where the concentration of the chemical dispensed by the supply and ratio of the flow rate of the injected chemical to the flow rate of the ozone-gas-water solution through the injector/mixer determines the concentration of the injected chemical in the solution exiting from the mixer. The supply 42 may be implement with a source of pressurized nitrogen regulated to a predetermined pressure (not shown) connected through a length of tubing to a reservoir (not shown) containing a liquid chemical to be injected. The dip-tube outlet (not shown) of the chemical reservoir is connected through a length of tubing to the inlet side a flow controlling needle valve (not shown). The outlet of the flow controlling needle valve is connected through a length of tubing to the inlet of a flow meter. The outlet of the flow meter is connected through a length of tubing to the inlet of the chemical injection control valve 44. The outlet of the chemical injection control valve 44 is connected through a length of tubing to the chemical injection port of chemical



injector/mixer 40. The pressurized injected chemical supply can also be implemented using a metering pump of other means known to those skilled in the art.

Point-of-use Chemical Injector: The chemical injector/mixer 40 may be a venturi injector, a static mixer, a mixing "T", or other device known to those skilled in the art. The injected chemical supply must deliver the chemical to the injection port of chemical injector/mixer with sufficient pressure to achieve the desired predetermined injected chemical flow rate. The internal volume of the injector mixer 40 must be kept to minimize the additional volume between the heat exchanger 28 and the dispense nozzle 36 referred to in the earlier discussion.

Minimizing temperature or Dissolved Ozone Concentration Changes Caused by the Injection of Chemicals: In the preferred embodiment the chemical injector/mixer 40 is located downstream of the heat exchanger 28. The temperature of the ozone-water solution is set at a predetermined temperature which is in the range of 30 to 95 deg. C. In the electronic device cleaning and processing embodiment the temperature may be approximately 50 deg. C. If the temperature of the injected chemical is below the temperature of the ozone-water solution entering the injector, then the solution exiting from the heater will be below the temperature of the solution entering the point-of-use heater. The fall in temperature may be mitigated by minimizing the volume flow rate of the injected chemical relative to the volume flow rate of the ozone-water solution. In an alternative embodiment, the injected chemicals can be preheated to approximately the same temperature as the temperature of the ozone-water solution entering the chemical inject thereby eliminating any fall in temperature mentioned above.

It is well known that by the introduction of hydroxyl radical scavengers (carbonates, bicarbonates, phosphates, etc) into an ozone-water solution at a molar concentration of 5 to 10 times the molar concentration of the ozone, that the concentration of the molecular ozone in solution can be maintained. We have found that borates can also be used to stabilize the concentration and adjust the pH. In some applications such as medical instrument sterilization, where metal ion contamination is not a concern, one can use a large range of hydroxyl radical scavenger chemicals such a sodium phosphate, potassium phosphate, sodium carbonate. In the case of electronic device processing, one can use ammonium counter ions in lieu of the metal counter ions. However, many candidate hydroxyl radical scavenger chemicals such ammonium carbonate, ammonium bicarbonate, ammonium phosphate, ammonium acetate, carboxylic acid, phosphonic acid, and salts thereof, as well as sulfates, for example ammonium sulfate, if given sufficient time, may react with ozone-water solution,

oxidize the injected chemical, consume ozone, and thereby significantly reduce the dissolved ozone concentration. The advantage of injecting these chemicals near the point-of-use is that the time available for these reactions to proceed very far is severely reduced. Accordingly, many chemicals which would not normally be useable if mixed in a heated ozone-water solution and given time to react can be employed by injecting the chemicals into the ozone-water solution stream just upstream of the point-of-use.

In the semiconductor wafer processing application a chemical may be injected at approximately 20-25 deg. C, at a flow rate of approximately 1 ml/sec into an ozone-water solution flowing at approximately 50 ml/sec. We found it useful to add approximately 1 ml of 1.0 mole/liter solution of a hydroxyl radical scavenger such as ammonium bicarbonate for every 50 ml of a dispensed 72 mg/Liter ozone-water solution (50:1 dilution of the injected mixture), thereby forming a mixture of ozone-water solution with a dissolved ozone concentration of about 72mg/Liter (1.5 millimoles/liter) and a hydroxyl radical scavenger concentration of about 20 millimoles/liter. Other alternative hydroxyl radical scavenger chemicals well known to those skilled in the art such as ammonium carbonate, ammonium acetate, and acetic acid can also produce good results

The pH of the ozone-water solution has a number important effects. First, the pH can influence metal corrosion rates. Generally, the optimum pH for minimizing metal corrosion is slightly less than 7. Second, the pH can influence of etch rate. For example, the inventors have shown that normalized etch rate of Shipley UV-6 DUV positive photoresist at pH=4.2 is about 1/10<sup>th</sup> the etch rate at PH=6.7. Many metal free pH adjusting chemicals suitable for electronic device manufacturing (ammonium hydroxide, ammonium phosphate monobasic, ammonium phosphate dibasic), if given sufficient time, may react with ozone-water solution, oxidize the injected chemical, consume ozone, and thereby significantly reduce the dissolved ozone. Point-of-use injection of these chemicals can significantly reduce the amount of consumption of both ozone and the chemical.

Metal corrosion control is a challenge in many aqueous cleaning systems. Corrosion inhibitors such as benzotriazoles, tolytriazoles, mercaptobenzathiazol, axoles, imidazoles, thioxoles, indoles, pyrazoles, benzoate, molybdates, phosphates, chromates, dichromates, tungstate, silicates, vandate, and borate may be introduced into the water-ozone solution to control metal corrosion. Benzotriazole is an attractive copper corrosion inhibitor. Corrosion inhibitor chemicals may be conveniently injected and mixed into the ozone-water solution just upstream of the point of application of the solution to the material to be cleaned or oxidized.

Surfactants are often used in aqueous cleaning systems to improve wetting of surfaces. However, most candidate surfactants react with ozone. Point-of-use injection of these chemicals can significantly reduce the amount of consumption of both ozone and the surfactant.

In the preferred embodiment the chemical injector/mixer 40 is located downstream of the heat exchanger 28. Alternatively, the chemical injector/mixer 40 could be located just upstream of the heat exchanger 28. In this case the injected chemicals are heated with the ozone-water solution. However, the chemicals have a slightly longer time available to react with the ozone-water solution.

#### Materials Processing Method Additionally Including Point-Of-Use Chemical Injection - Multiple Chemical Supplies

Description- Fig. 4

In an alternative embodiment, means may be provide for the injection of more than one chemical, each from a separate chemical reservoir. With reference to Fig 4, an ozonated water supply 22 is connected through a length of tubing to the common input of a three-way valve 24. The one outlet of three-way valve 24 is connected through a length of tubing to the cold process fluid inlet of a heat exchanger 28. The other outlet of three-way valve 24 is connected through a length of tubing to the facility drain-reclaim 26 for the ozone-water solution. The heated working fluid outlet of a recirculating heating unit 30 is connected through a length of tubing to the heated working fluid inlet of heat exchanger 28. The heated working fluid outlet of heat exchanger 28 is connected through a length of tubing to the working fluid return of recirculating heating unit 30. The heated process fluid outlet of heat exchanger 28 is connected through a short length of tubing to the inlet of chemical injector/mixer 41. The outlet of an injected chemical supply 40-1 is connected to the inlet of two-way valve 44-1. The outlet of valve 44-1 is connected to a first chemical injection port of a multiple port chemical injector/mixer 41. The outlet of an injected chemical supply 40-2 is connected to the inlet of two-way valve 44-2. The outlet of valve 44-2 is connected to a second chemical injection port of a multiple port chemical injector/mixer 41. The outlet of an injected chemical supply 40-3 is connected to the inlet of two-way valve 44-3. The outlet of valve 44-3 is connected to a third chemical injection port of a multiple port chemical injector/mixer 41. The outlet of chemical injector/mixer 41 is connected to the common inlet port of three-way valve 32. The one outlet of three-way valve 32 is connected through a short length of tubing to a dispense nozzle 36. The dispense nozzle 36 is spaced a relatively

short distance from the surface 38 of the material to be oxidized, cleaned, or processed. The other outlet of three-way valve 32 is connected through a length of tubing to the facility drain-reclaim 34 for the heated ozone-water solution. Embodiments for the injection of chemicals, either from a fewer number, or from a greater number of injected chemical supplies, can be implemented using a similar approach.

Separate Injector/Mixer Element for Each Injected Chemical. In an alternative embodiment a separate injector/mixer element for each injected chemical, may be used in lieu of a single injector/mixer element, with multiple chemical inlet ports. The design of apparatus for the injection of chemicals into a fluid stream is well known to those skilled in the art. Many alternative approaches may be chosen provided that the residence time of the ozone-water solution in the injecting mixing element(s) is small since this residence time adds to the total residence time. The short residence time from the inlet of the point-of-use heater 28 or 29 to the point of application of the ozone-water and injected chemical solution to the material to be oxidized has two important benefits. A short residence time minimizes the amount of time available for the dissolved ozone concentration of the supersaturated ozone-water solution entering the mixing element to fall much during the time required for the solution to pass through the element. A short residence time also minimizes the amount of time available for the ozone-water solution to react with the injected chemicals. If the chemicals react with the ozone-water solution, the reaction may not only consume ozone and reduce the dissolved ozone concentration, but also may consume some or all of the injected chemical.

Means for Purging Chemical Injector Lines. In an alternative embodiment, each chemical injection valve 44-1, 44-2, and 44-3 may be a four-way valve to provide for purging the injection line. In the "off" position a four-way chemical injection valve can shut off chemical injection to the injected chemical inlet. In the "on" position, the valve can permit chemical injection to the injected chemical inlet. In the "purge" position permit the purging with DI water, for example, the lengths of tubing between the valves 44-1, 44-2, and 44-3 and the chemical injector/mixer 41 to prepare for the introduction of a different chemical into the injected chemical inlet of chemical injector/mixer 41.

#### Operation - Fig. 4

This embodiment may provided for the injection of different chemicals a predetermined times during the materials processing cycle. The rate of injection for each of the chemicals can be specified and controlled for each instant of time during the materials processing cycle. A preferred embodiment may utilize a computer or microprocessor to control the flow rates at each time step of the process. The operation of the of the point-of-

use chemical injection system with multiple chemical supplies is otherwise the same as the operation of a point-of-use chemical injection system with a single chemical supply.

### Materials Processing Method With A Spin Processor

#### Description - Fig. 5

One preferred method for applying the ozone-water solution to semiconductor substrates and the like is to apply the ozone-water solution to the surface of the substrate while spinning the substrate about an axis at a relatively high (1,000 to 4,000 rpm) rotational speed. The use of a this method for applying the ozone-water solution to semiconductor substrates and the like provides for a higher mass transport rate  $M$  than can be achieved by immersion techniques. With reference to Fig 5, an ozonated water supply 22 is connected through a length of tubing to the common input of a three-way valve 24. The one outlet of three-way valve 24 is connected through a length of tubing to the cold process fluid inlet of a heat exchanger 28. The other outlet of three-way valve 24 is connected through a length of tubing to the facility drain-reclaim 26 for the ozone-water solution. The heated working fluid outlet of a recirculating heating unit 30 is connected through a length of tubing to the heated working fluid inlet of heat exchanger 28. The heated working fluid outlet of heat exchanger 28 is connected through a length of tubing to the working fluid return of recirculating heating unit 30. The heated process fluid outlet of heat exchanger 28 is connected through a short length of tubing to the inlet of chemical injector/mixer 40. The outlet of an injected chemical supply 42 is connected to the inlet of two-way valve 44. The outlet of valve 44 is connected to the chemical injection port of chemical injector/mixer 40. The outlet of chemical injector/mixer 40 is connected to the common inlet port of three-way valve 32. The one outlet of three-way valve 32 is connected through a short length to one inlet of a three-way valve etch/rinse valve 46. The common outlet of three-way etch/rinse valve 46 is connected through a short length of tubing to the process fluid inlet 48 to a dispense nozzle 36 located in a gas tight materials processing module 50. In the preferred embodiment the materials processing module 50 is fitted with a lid with a gas tight seal (not shown). The sealed materials processing module serves to contain the any ozone gas that is released from solution at the point of application. A pressurized DI water rinse supply 74 is connected through a length of tubing to the other inlet of port of the three-way etch/rinse valve 46. Pressurized DI water rinse supply 74 typically comprises a pressurized source of DI water connected though a liquid pressure regulator and a liquid flow controller and liquid particulate filter. The dispense nozzle 36 is spaced a relatively short distance (typically 0.5 to 10 cm) from the

surface of the material 38 to be oxidized, cleaned, etch, or processed. The other outlet of three-way valve 32 is connected through a length of tubing to the facility drain-reclaim 34 for the heated ozone-water solution.

A process fluid drain outlet port 52 of the materials processing module 50 is connected through a short length of tubing to the inlet of liquid trap 54. When the ozone-water solution is dispensed onto the surface of the substrate at a given flow rate, the gas displaced from the housing may exit at approximately the same flow rate with a relatively small pressure drop. This insures that the pressure in the housing does not rise during the dispense cycle. The outlet of liquid trap 54 is connected by a length of tubing to the facility drain/reclaim 56 for process module liquid effluent. The wafer or substrate or the like is held by a spinner chuck 58. The chuck 58 may hold the wafer or substrate or the like (the material 38 to be oxidized, cleaned, etched, or processed) by the use of vacuum, edge clamps, or other means well known to those skilled in the art. The spinner chuck 58 is connected by a shaft or other means to wafer spinner motor 60. The motor, typically controlled by a microprocessor, can be programmed to accelerate the wafer or substrate or the like at a predetermined rate from zero rpm to a predetermined rpm, hold that rpm for a specified period of time, then decelerate at a predetermined rate to back to zero. Alternatively, the motor may be programmed to successively spin at several different rpm values chosen for different portions of the materials processing cycle. An ozone off-gas outlet port 62 is connected through a length of tubing to the inlet to an ozone catalytic unit 64. The outlet of ozone catalytic unit 64 is connected through a length of tubing to a facility exhaust vent 66. The diameter of the vent line and flow capacity of the ozone catalytic unit 64 is chosen so that the vent line is non-back pressuring. In the preferred embodiment the ozone destruction unit 64 is of the catalytic type filled with a catalyst such as Carulyte 200 (Carus Corporation). Since the waste ozone gas entering the catalyst unit contains water vapor, one may heat the catalyst unit with a heat tape and temperature controller to about 50 degree C to prevent moisture condensation on the catalyst. The increased temperature also serves to increase the performance of the catalyst. The catalyst unit is sized to provide a sufficient residence time to convert the high concentration waste ozone off-gas to oxygen. The higher the waste gas flow rate, the larger the catalyst volume must be to achieve the conversion. The required residence time can be obtained from the catalyst supplier. Alternatively, the ozone destruction unit 64 can be thermal destruction unit in which the ozone gas-oxygen mixture is decomposed back to oxygen by raising the temperature of the waste ozone gas to about 300 degree C. A pressurized nitrogen purge gas supply 68 is connected through a length of tubing to the inlet

port of a two-way valve purge valve 70. The outlet port of the two-way gas purge valve 70 is connected to a purge-gas inlet 72 to the materials processing module 50. Pressurized nitrogen purge supply 68 typically comprises a pressurized source of nitrogen connected through a gas pressure regulator and a gas flow controller and gas particulate filter.

In a single wafer processing embodiment one can use a single wafer spinner such as the all Teflon microprocessor controlled wafer spinner made by Laurell in which the spinner has been fitted with a gas tight lid seal. The Laurell spinner model WS400-6TFM/Lite is all Teflon and designed to accommodate wafers up to 150 mm diameter. The spinner acceleration rate, deceleration rate, and rpm can be set and controlled between 100 RPM and 6000 RPM. The spinner can be upgraded with a valve control option that enables the spinner microprocessor to control up to eight valves. This enables precise automated control of the purge to waste, etch dispense, and DI rinse dispense cycle times. Laurell makes spinners to accommodate 200 mm diameter wafers, 300 mm diameter wafers, and other substrates sizes and shapes.

#### Measured Photoresist Removal Performance with Equilibrium Processing with $T_2=T_1$

We have measured photoresist etch rate using a spin-etch configuration where ozone gas was dissolved in water at a given temperature  $T_1$  and given concentration. The ozone-water solution was dispensed at the same temperature  $T_2$  through a dispense nozzle to the center of spinning wafer coated with a layer of hard baked I-line positive photoresist or DUV positive photoresist.

We used 150 mm diameter silicon wafers coated with a thin (0.5 to 0.8 micron) and uniform layer of photoresist. We exposed the layer to a stream of ozone dissolved in water at a given concentration and temperature for a specified period time to partially remove some of the thickness of resist. We then measured the remaining layer thickness with an NanoSpec ellipsometer and calculated an etch rate, the amount of organic material removed per unit time. The results are summarized in Tables 7 and 8 below.

**Table 7** The measured etch rate when ozone is dissolved in water at a given temperature and given concentration and applied at a given flow rate through a dispense nozzle at the same temperature to the center of spinning wafer coated with a layer of hard baked I-line positive photoresist. The highest etch rate is achieved at the center of the wafer and the lowest etch rate is achieved at the edge of the wafer for the example shown.

Dissolved Ozone Concentration (mg / L)	Dissolved Temp (deg C)	Applied Temp (deg C)	Flow Rate (L/min)	Wafer spin (RPM)	Max Etch Rate (Å°/min)	Min Etch Rate (Å°/min)
75	9.9	9.9	1.2	4000	4270	3718
62	18.8	18.8	1.26	4000	6178	3538
24.9	27.8	27.8	1.26	4000	3524	1940

**Table 8** The measured etch rate when ozone is dissolved in water at a given temperature and given concentration and applied at a given flow rate through a dispense nozzle at the same temperature to the center of spinning wafer coated with a layer of hard baked DUV positive photoresist. The highest etch rate is achieved at the center of the wafer and the lowest etch rate is achieved at the edge of the wafer for the example shown.

Dissolved Ozone Concentration (mg / L)	Dissolved Temp (deg C)	Applied Temp (deg C)	Flow Rate (L/min)	Wafer spin (RPM)	Max Etch Rate (Å°/min)	Min Etch Rate (Å°/min)
90	10.4	10.4	1.2	4000	1226	986
60	19.2	19.2	1.32	4000	2426	1774
24.9	27.8	27.8	1.23	4000	2308	1058

We know from our previous discussion that increasing the temperature of the water reduces the equilibrium dissolved ozone concentration as predicted by Henry's law. In the case of semiconductor wafers coated with a layer of organic polymer, we found that etch rate was higher at 20 degree C than at 10 degree C even though the equilibrium dissolved ozone concentration was significantly lower (60 mg/L at 20 deg. C compared to 90 mg/L at 10 deg. C). We see that the maximum etch rate first increases with increased temperature and then the



etch rate decreases with further increases in temperature. We see that I-line positive photoresist is etched by the process at a faster rate than DUV resist. This is because the I-line positive photoresist has a higher value of S than DUV positive photoresist at a given temperature.

These etch rates are not high enough to provide reasonable throughputs in a single wafer configuration.

#### Measured Photoresist Removal Performance with Non-Equilibrium Processing with $T_2 > T_1$

We tested one embodiment of the invention in which which ozone is dissolved in water at a lower temperature to produce a high dissolved concentration, then passed at given flow rate though a point of use heater, and applied at a higher temperature to the center of spinning wafer. with both I-line and DUV positive photo resist and showed that we were able to achieve an I-line positive photoresist etch rate of 16,812 A°/minute and DUV positive photoresist etch rate of 10,626A°/minute. (See Tables 9 and 10). These etch rates are about a factor of five higher than the highest etch rates than we were able to initially achieve using conventional techniques (compare to Tables 7 and 8). The mass transport M is generally higher at a higher RPM and the etch rate is higher as well.

We again see that I-line positive photoresist is etched by the process at a faster rate than DUV resist because the I-line positive photoresist has a higher value of S than DUV positive photoresist at a given temperature.

Table 9 The measured etch rate when ozone is dissolved in water at a lower temperature to produce a high dissolved concentration, then passed at given flow rate though a point of use heater, and applied at a higher temperature to the center of spinning wafer. The wafer is coated with a layer of hard baked I-line positive photoresist. The highest etch rate is achieved at the center of the wafer and the lowest etch rate is achieved at the edge of the wafer.

Dissolved Ozone Concentration (mg / L)	Dissolved Temp. (deg C)	Applied Temp. (deg C)	Flow Rate (L/min)	Wafer spin (RPM)	Max Etch Rate (A°/min)	Min Etch Rate (A°/min)
95	7.5	49.5	2.7	1000	20208	9459
95	7.5	47.5	2.7	4000	23850	16812

Table 10 The measured etch rate when ozone is dissolved in water at a given temperature and given concentration and applied at a given flow rate through a point of use heater and through a dispense nozzle at a higher temperature to the center of spinning wafer coated with a layer of hard baked DUV positive photoresist. The highest etch rate is achieved at the center of the wafer and the lowest etch rate is achieved at the edge of the wafer.

Dissolved Ozone Concentration (mg / L)	Dissolve Temp. (deg C)	Applied Temp. (deg C)	Flow Rate (L/min)	Wafer spin (RPM)	Max Etch Rate (Å°/min)	Min Etch Rate (Å°/min)
95	7.5	48.5	2.7	1000	16227	6117
95	7.5	47.5	2.7	4000	19413	10626

We have presented experimental results on a method for removing photoresist at high speed using a solution of ozone gas dissolved in water comprising the steps of a) dissolving high ozone gas in water at a relatively low predetermined temperature to form an ozone-water solution with a high dissolved ozone concentration, b) heating the ozone-water solution with a point-of-use water heater (a liquid to liquid heat exchanger) to quickly increase the solution temperature to a predetermined higher temperature, and c) applying the heated ozone-water solution to the center of a photoresist coated wafer rotating at high speed. The etch rate is increased, not only because the surface reaction rate is increased at the higher temperature, but also because the heated ozone-water solution has a much higher dissolved ozone concentration at the higher temperature than could be achieved if the ozone gas was initially dissolved in water at the higher temperature under equilibrium conditions

Alternative Means for Applying Ozone-Water Solution to The Substrate: The application of the ozone-water-other chemicals solution to the surface or surfaces of the material to be processed can be accomplished in a number of different ways. In the preferred embodiment the solution can be applied to the center of the wafer through a single solid stream nozzle 36 with an inside diameter of about 6 mm positioned to apply a flow of water to the wafer at the center. In a second embodiment the solution can successively applied to different positions between the center and edge of the wafer. In this second embodiment the ozone-water solution can be flowed through a nozzle which can be successively positioned at different locations from the center to the edge of the wafer or from the edge to the center of the wafer. The dwell time at each position can be controlled to reduce the radial variation in

the etch rate or cleaning rate or oxidation rate over the duration of the materials processing cycle. In a third embodiment the solution can be applied the surface of the wafer with multiple nozzles 36A, 36B, 36C, ... (not shown) In the fourth embodiment one or more nozzles may be mounted on one or more rotating spray arms (not shown) positioned to apply the solution to one or more surfaces of the material to be oxidized. The ozone-water-other chemicals solution can be applied to the surface or surfaces of the material to be processed by other means familiar to those skilled in the art.

Alternative Means for Applying DI Rinse Water to The Substrate. In an alternative embodiment the DI rinse water may be applied to the substrate with a separate set of one or more rinse nozzles (not shown). Rinse nozzles may be chosen for optimum rinse performance at a predetermined rinse flow rate and the nozzles may be positioned to rinse one more surfaces of the substrate.

Optional Instrumentation Upstream of the Dispense Point: A dissolved ozone monitor and temperature sensor may be inserted in the short length of tubing just upstream of the dispense nozzle 36. This instrumentation provides a continuous readout of the dissolved ozone concentration and temperature of the heated ozone-water solution just upstream of the dispense point. This can be a source of useful diagnostic information during process development. Since the elements add additional volume between the inlet of the water heater 28 and the point of application of the solution to the material 38, the internal volume of these optional instrumentation sensors should be small so that the residence volume through which the heated ozone-water solution must pass is kept small and the time delay between heating the ozone-water solution and applying the heated ozone-water solution to the material 38 is kept sufficiently small as discussed earlier.

Alternative Locations of the Back Pressure Regulator for an Ozonated Water Supply Designed to Dissolve Ozone Gas in Water at a Pressure Greater Than 1 Atmosphere (14.5 psia.) In an ozonated water supply designed to dissolve ozone gas in water at a pressure above 14.5 psia, a gas back pressure regulator is typically placed in the waste-ozone off-gas (undissolved ozone gas) outlet of the ozone gas-water contactor element and a liquid back pressure regulator is typically placed in the ozonated outlet line of the ozone-gas water contactor element. In the preferred embodiment for spin processing or semiconductor wafers and the like, the apparatus is designed to apply the ozone gas-water solution to the material to be oxidized 38 in a material processing module operated at one atmosphere pressure. In one embodiment the liquid back pressure regulator may be positioned between the outlet of the ozonated water supply 22 and the inlet of the two three-way valve 24. Alternatively, the

liquid back pressure regulator may be moved to a position just upstream of the dispense nozzle 36 to minimize the time available for the ozone gas to leave the ozone-water solution from the time the pressure is reduced to atmospheric pressure and the solution is applied to the material. In systems designed to apply the ozone gas-water solution to the material to be oxidized 38 in a material processing module operated at the same pressure as a pressurized ozone-gas-water contactor element, the liquid back pressure regulator may be positioned at the liquid outlet of a pressurized materials processing module and the gas back pressure regulator, which is typically placed in the waste-ozone off-gas outlet of the ozone gas-water contactor, can be relocated to the waste-ozone off-gas outlet of the materials processing module.

#### Operation - Fig. 5

A preferred technique for applying the ozone-water solution to semiconductor substrates and the like is to apply the ozone-water solution to the surface of the substrate while spinning the substrate about an axis at a relatively high (1,000 to 4,000 rpm) rotational speed. The use of this method for applying the ozone-water solution to semiconductor substrates and the like provides for a higher mass transport rate  $M$  than can be achieved by immersion techniques. One very important use of the preferred embodiments is for the removal of photoresist and post etch residue from semiconductor wafers and the like. Let us describe the operation for a typical photoresist or post etch residue removal application. With reference to Fig 5, ozonated water supply 22 supplies an ozonated water formed by dissolving ozone gas at a gas phase concentration of 240 mg/L and pressure of 14.5 psia (1 bar) into DI water chilled to a temperature of about 8 degree C. The ozonated water supply delivers the chilled ozone gas-water solution at a dissolved concentration of about 90 mg/L and at a flow rate of 2.7 Liter/min through three way purge valve 24, through heat exchanger 28 or point-of-use heater 29 where the solution temperature is increased to about 50 degree C, through chemical injector/mixer 40, through three-way purge valve 32, through three-way etch/rinse valve 46, through materials-processing-module inlet 48, through dispense nozzle 36 where the heated ozone-water solution is applied to the center of a semiconductor wafer 38 spinning at about 3500 to 4,000 rpm. Under these conditions the inventors have shown that the dissolved ozone concentration downstream of the point-of-use heater is approximately 75 mg/L, more than 80 percent of the concentration at the inlet of the point-of-use heater. The ozone-water solution traverses the surface to the wafer from the point of application to the edge of the wafer and enters the process fluid outlet of the materials-processing-module where the ozone-water solution and other liquid effluents from the process are carried through a trap

to facility drain 56. The trap prevents back flow of an gases from the facility drain reclaim reservoir. The nitrogen purge supply flows dry filtered nitrogen gas, or another suitable gas, at a flow rate of about 0.5 L/min, through the two-way nitrogen purge valve 70, to the inlet of the materials-processing-module 72. The nitrogen gas assists in the removal of any ozone gas that leaves the ozone-water solution inside the materials-processing-module and provides an nitrogen blanketed processing environment. The ozone off-gas, nitrogen, and other waste gases from the process, exit from the materials-processing-module at outlet port 62, pass through the catalyst unit 64 where the waste ozone gas is converted to oxygen before exiting to the facility exhaust vent 66. The period during which the ozone-water solution is applied to the spinning substrate can be designated as the duration of the etch clean cycle. Once the etch cycle is completed, the three way purge valve 32 can be set to direct the flow of the heated ozone-water solution to the facility drain/reclaim 34 for the heated ozone-water solution and the three-way etch/rinse valve 46 can be placed in the rinse position to allow the rinse water to flow to the materials-processing-module. DI water can then flow from the pressurized DI water supply through the three way valve 46, through materials-processing-module inlet 48, through dispense nozzle 36 where the DI water solution is applied to the center of a semiconductor wafer 38 spinning at about 3500 to 4,000 rpm. The period during which the DI rinse water is applied to the spinning substrate 38 can be designated as the duration of the rinse cycle. Once the rinse cycle is completed, the three-way etch rinse valve 46 can be returned to the etch position while the three-way purge valve 32 remains set to direct the flow of the heated ozone-water solution to the facility drain/reclaim 34 so that all liquid flows to materials-processing-module are off and the wafer can be spun dry. The period during which all liquid flows to the materials processing module are off and the substrate is spinning can be designated as the duration of the spin dry cycle. The spin RPM and duration of each cycle can be set to a predetermined value for a particular process application.

Example Process Conditions for Photoresist Removal And Post Etch Residue Removal. A typical wafer spin processing sequence may include spin etch or cleaning or oxidation cycle, a spin rinse cycle, and a spin dry cycle. Example process conditions for photoresist removal and post etch residue removal with 150 mm diameter wafers in a single wafer spin processing configuration are summarized in Table 11 below.

Table 11 Example Process Conditions for Photoresist Removal And Post Etch Residue Removal- Single Wafer Processing Configuration - 150 mm Diameter Wafers -

<u>ETCH CLEAN</u>	
Dissolved Ozone Concentration (mg/L)/(millimoles/liter)	70 to 210 mg/L (1.5 to 4.5 millimoles/L)
Hydroxyl Radical Scavenger	Carbonate, Bicarbonate, Phosphate, Acetate
Hydroxy Radical Scavenger Concentration (millimoles/liter)	5 to 10 x dissolved ozone concentration
pH	6.5 to 7.5
Ozone-Water Solution Temperature at the point of application (°C)	40 to 60
Spin Etch Speed (RPM)	2,000 to 4,000
Ozone-Water Solution Flow Rate (L/min)	1.0 to 3.0
Etch Duration (min)	0.5 to 6.0
<u>RINSE</u>	
DI Rinse Temperature (°C)	20
Spin Rinse Speed (RPM)	2,000 to 4,000
DI Rinse Flow Rate (L/min)	0.5 to 1.0
DI Rinse Duration (min)	0.33 to 1.0
<u>DRY</u>	
Spin Dry Speed (RPM)	2,000 to 4,000
Dry Duration (min)	0.33 to 1.0

These process conditions are typical for this type of application. However, process conditions outside the range of values presented in the Table 11 above can also provide satisfactory performance for this and other applications and wafer processing configurations. For example, batch wafer spinning configurations in which two to four cassettes of wafers are processed at one time typically operate at a lower RPM and lower etch flow rate per wafer. The RPM for a batch spinner is typically in the range of 500 to 1500 RPM. The total etch chemistry flow rate for a batch wafer spinner is typically in the range of 10 to 20 liters/minute.

The lower RPM and lower flow rates will yield a lower mass transport rate and lower etch rate. The temperature at which the etch rate will become mass transport limited will be lower since the mass transport rate is lower as discussed earlier.

Process Performance Measured by the Inventors: A typical duration for an etch cleaning cycle for removing photoresist and post etch residue is 0.5 to 6 minutes. The inventors have demonstrated the ability to remove hard baked Olin OCG-897 I-line positive photoresist from semiconductor wafers at an etch rate of approximately 17,000 Angstroms/minute. The inventors have demonstrated the ability to remove hard baked Shipley UV6 DUV positive photoresist from semiconductor wafers at an etch rate of approximately 11,000 Angstroms/minute. The etch rate is generally lowest at the wafer edge. The reported etch rates are those measured at the wafer edge at a flow rate of 2.7 L/min., a dissolved ozone concentration of about 75 mg/L, a spin speed of about 4,000 RPM, and temperature of approximately 50 degree C, and hydroxyl radical scavenger concentration of about 15 millimoles/Liter. The inventors have demonstrated the ability to remove both post metal etch resist and resist residue under similar conditions from test structures using 0.35  $\mu\text{m}$  technology with I-line resist in less than 1.5 minutes at 50 degree C. The inventors have demonstrated the ability to remove both post metal etch resist and resist residue from test structures using 0.18  $\mu\text{m}$  technology with DUV resist in less than 3.0 minutes at 50 degree C. It is useful to compare to cycle times of this new ozone-water post etch cleaning process with the current process based upon plasma ashing and solvent residue removal. This is summarized in Table 12 below.

		CONVENTIONAL PROCESS		NEW OZONE-WATER PROCESS		
		I-line resist	DUV resist		I-line resist	DUV resist
Step #	Process Step	Time Req'd (min)	Time Req'd (min)	Process Step	Time Req'd (min)	Time Req'd (min)
	single wafer process			single wafer process		
1	O2 Ash	3.0	5.0	Ozone Etch	1.5	3.0
2	Hot/Cold DI Rinse	1.5	1.5	Cold DI Rinse	0.5	0.5
3	Spin Dry	0.5	0.5	Spin Dry	0.5	0.5
	batch process					
4	Batch Solvent Clean	10	10	NA		
5	Batch DI Rinse	5	5	NA		
6	Batch Spin Dry	5	5	NA		

The process has demonstrated the ability to remove the post oxide etch resist and resist residue from the test structures using 0.35  $\mu\text{m}$  technology with OCG 897-12 positive I-line



resist in 1.0 minutes. The process has demonstrated the ability to remove the post oxide etch resist from the test structures using 0.18  $\mu\text{m}$  technology with Shipley UV6 positive DUV resist in 3.0 minutes. However, the process has not yet demonstrated the ability to completely remove the post-oxide-etch residue from the test structures using 0.18  $\mu\text{m}$  technology with Shipley UV6 positive DUV resist with a cycle time of 5 minutes. In applications using I-line resist only the new ozone-water process has the potential to replace the conventional plasma ashing and solvent cleaning process used for post-oxide etch clean.

#### Materials Processing Method With Spray Processor

##### Description - Fig. 6

There are a number of applications in which one may wish to process materials such as medical instruments, bio-materials, and medical devices for the purpose of surface treatment, cleaning, disinfection, or sterilization. In these types of applications alternative processing configurations can be used. The principal difference between the spray processor and the spin processor shown in Fig. 6 is that the spray processor does not spin the material to be processed at high RPM. The materials to be processed may not be able to be spun at high RPM because of their mass, size, or asymmetric shape. In this case the instruments or materials to be processed can be placed on a rack or wire mesh support and the heated ozone-water-chemical solution can be applied to the materials from multiple directions using one or more spray heads. The materials may be mounted on a rack or supporting structure that permits good access for the spray and the heated ozone-water-chemical solution may be applied with one or more spray heads. The spray heads may be fixed in position, mounted on rotating spray arms, mounted on translating spray arms. The rack or structure that supports the materials to be processed may be fixed in position, may be slowly rotated at low rpm, or may be slowly translated. A materials processing module designed for sterilization of materials or devices or instruments must have several additional elements to prevent microbial contamination of the materials processing module. The additional elements may be understood by referring to Fig. 6. The elements 22, 24, 26, 28, 29, 30, 32, 34, 36, 38, 40, 42, 44, 48, 50, 52, 54, 56, 62, 64, 66, 68, 70, and 72 are the same as defined earlier. A materials processing module designed for sterilization does not require a separate rinse supply. Accordingly, the rinse supply 74 and etch rinse valve 46 is not required. Three-way valve 32 is connected by short lengths of tubing through the process fluid inlet 48T and 48B to top spray assembly 36T and bottom spray assembly 36B located above and below a support structure 76. A drain line check valve 78 is located between the process-fluid outlet 52 and the trap 54 to prevent backflow to the process

module. A vent line check valve 80 is located between the off-gas outlet 62 and the catalytic ozone destruction unit 64 to prevent backflow to the process module. A purge gas sterile filter 82 is located between the outlet of two-way purge gas valve 70 and the purge gas inlet 72.

#### Operation - Fig. 6

In reference to Fig. 6, the operation of the spray processor is similar to that of the spin processor. The ozone gas-water solution is formed at a relatively low temperature and heated at the point-of-use as described earlier. Additional chemicals can be injected at the point-of-use from one or more chemical supplies as described earlier. The heated ozone-water solution flows from the outlet of three-way valve 32 through the process fluid inlets 48T and 48B in the spray type materials processing module 50-S to the top and bottom spray assemblies 36T and 36B. The material to be processed or sterilized 38 is positioned on an open support structure 76. The support structure is designed to not block the spray from impinging on the surfaces of the material to be processed or sterilized. The heated ozone gas-water solution is sprayed onto all surfaces of the materials or instruments or devices with the top and bottom spray assemblies 36T and 36B. If the instruments have internal channels that must be sterilized, then the heated ozone gas -water solution may be also flowed through those internal channels as well. Ozone dissolved in water is a zero residue sterilant. However, if additional chemicals such as surfactants are injected and mixed with ozone-water solution to improve wetting and soil removal during the sterilization cycle, then the residues of those added chemicals may need to be rinsed from the surfaces of those instruments or materials in a subsequent sterile rinse cycle.

A sterile water rinse method not requiring a sterile filter. In the prior art the sterile rinse has been implemented by passing water through a 0.2 micron sterile filter. This type of filter becomes clogged with use and must be replaced at regular intervals. In the present embodiment, one can use the heated ozone gas-water solution as a sterile rinse solution. The solution may be used with no additional injected chemicals except perhaps a chemically benign buffer and scavenger. One good candidate buffer is a phosphate buffer made by mixing  $\text{KH}_2\text{PO}_4$  and  $\text{Na}_2\text{HPO}_4$ . Another candidate buffer is boric acid and sodium borate. The phosphate or borate will also serve as a hydroxyl radical scavenger. The ozone gas dissolved in water to form a supply of ozonated water will quickly kill the most resistant organisms. The 12D sterilization time for the most resistant water-borne organisms such as *Giardia* and *Cryptosporidium* at a dissolved ozone concentration of only 60 mg/L is 0.2 minutes (12 seconds) as shown in Table 13. The residence time of a typical ozone gas-water

contactor used in an ozonated water supply is much longer than 12 seconds. The residence time in a recirculating contactor can be higher. Accordingly, the ozone-water solution dispensed from the ozonated water supply at a concentration of about 60 mg/L will be sterile water. When the ozone-water solution is passed through the point-of-use heater 28 or 29 and heated to a higher temperature (50 degree C for example), the concentration will decay to a low value with an exponential decay time constant estimated by the inventors to be about 25 seconds. The materials processing chamber may be continuously purged with sterile nitrogen or sterile air during the sterilization cycle to sweep the waste ozone off-gas from the chamber. When the instruments or devices are removed from the instrument chamber at the end of the sterilization and optional rinse cycle, the dissolved ozone concentration will have decayed to a very low value. Example process conditions for medical instrument and medical device sterilization are shown in Table 14.

Table 13: The lethality of ozone dissolved in water at a specified concentration against microorganisms in aqueous suspension in a stirred reactor. The time required to achieve a 12 log reduction in the specified organism at two different concentrations is calculated from the measured D value data. The D value is the time required to reduce the number of viable organisms by a factor of 10 at a specified sterilant concentration C

Microorganism	D	12CD	12D	note
	secs	mg.min/l	minutes	
	C=10 mg/l		C=60 mg/l	
Escherichia Coli	0.03	0.06	0.001	a
Striptococcus Fecalis	0.045	0.09	0.002	a
Mycobacterium Tuberculosis	0.15	0.3	0.005	a
Polio Virus	0.06	0.12	0.002	a
Endamoeba Histolytica	1	2	0.033	a
Bacillus Megatherium (Spores)	3	6	0.100	a
Giardia Lamblia, Giardia Muris (Cysts)	6	12	0.200	a
Cryptosporidium	6	12	0.200	a

Note a: Block, 1978; Sobsey, 1989: D values measured at a dissolved concentration of 1-2 mg/l, pH 7, and temperature of 15°C

Table 14: Example Process Conditions for Medical Instrument and Medical Device Sterilization

<u>STERILIZE</u>	
Dissolved Ozone Concentration (mg/L)/(millimoles/liter)	70 to 210 mg/L (1.5 to 4.5 millimoles/Liter)
Hydroxyl Radical Scavenger	Examples: Carbonate, Bicarbonate, Phosphate, Acetate
Hydroxyl Radical Scavenger Concentration (millimoles/liter)	5 to 10 x dissolved ozone concentration
pH Buffer	Examples: Phosphate, Borate
pH Buffer Concentration	~10 millimoles/Liter
pH	6.5 to 7.5
Surfactant - optional	Non-ionic and Anionic Mixture
Sequestering Agent - optional	Examples: Sodium Tripolyphosphate (STTP)
Sequestering Agent Concentration	10 millimoles/Liter
Ozone-Water Solution (Sterilant) Temperature at the point of application to the (°C)	30 to 50
Ozone-Water Solution Flow Rate (L/min)	5 to 10.0
Sterilization Cycle Duration (min)	5 to 10.0
<u>RINSE - optional</u>	
<u>DRY - optional</u>	

Process Performance Measured by the Inventors: The inventors have calculated the sterilant efficacy of an ozone-water solution at different solution temperatures for a dissolved ozone concentration of 60 mg/L based upon inactivation rates, measured by the inventors under equilibrium conditions, for AOAC porcelain penicylinders and AOAC Dacron polyester

suture loop carriers inoculated with IE6 *Bacillus Subtilis* spores according to the AOAC protocol.

Inactivation times are influenced not only by surface reaction rate but also by the dissolved concentration. They are also confounded by the statistical nature of the carrier inactivation process. We computed the average 12CD value for inactivation of carriers at each of the different equilibrium temperatures and dissolved concentrations. The use of the concentration-time product effectively normalized the results by the concentration. The results showed there is a very strong temperature dependence of inactivation rate. The analysis demonstrated that the inactivation at 20 deg. C is not mass transport limited, but surface reaction rate (temperature) limited.

The results for penicylinders are presented in Table 15. The parameter 12CD is the concentration-time product for inactivation of a million spore challenge with a million to one assurance level. We see that an increase in the temperature from 20 deg. C to 30 deg. C ( 10 deg. C increase), causes the 12CD value to decrease by more than a factor of 4 from 14 to 2.6. In next column (col. 6). We have computed the value of 12 CD at 10, 30, 40, and 50 deg. C from the measured value at 20 deg. C. The computation is based upon the following hypothesis: 1) the surface reaction rate coefficient S increases by a factor of 2 for every 5 deg. C increase in temperature (similar to that observed for the oxidation of polymer films), and 2) the overall reaction rate is dominated by the surface reaction rate over the temperature range. This assumption seems to be valid up to 30 deg. C. We have also computed the predicted 12D sterilization time at a dissolved ozone concentration of 60 mg/L (see table column 7). Further measurements will be required to determine the point at which the reaction becomes mass transport limited.

Table 15: The measured and calculated temperature dependence of the D value and 12 D sterilization time for porcelain penicylinders inoculated with bacillus subtilis (BS) var niger spores (ATCC 19659) according to AOAC protocol. The initial spore population was at least 1E6 CFU per carrier

Temp. deg. C	No. of Replicate Tests	Total No. of Carriers Tested	D value min. @ 60 mg/L	12CD hr. mg/L measured	12CD hr. mg/L <u>calculated</u> <u>from</u> <u>values @</u> <u>20 deg. C</u>	12D time minutes @ C = 60 mg/L
10					56	56
20	24	153	1.2	14	14	14
30	7	43	.3	2.6	3.5	3.5
40					.875	.875
50					.215	.215

The results for polyester suture loops are presented in Table 16. We see that polyester suture loops are more resistant to inactivation than penicylinders, and hence the FDA approved cycle time is generally determined by suture loop inactivation time. We see that an increase in the temperature from 10 deg. C to 20 deg. C to 30 deg. C ( 10 deg. steps), causes the 12CD value to decrease by nearly a factor of 4 for each step from 285 to 71 to 26. In next column(col. 6) we have again computed the value of 12 CD at 10, 30, 40, and 50 deg. C from the measured value at 20 deg. C based upon the assumption stated above. Again, the results are remarkably consistent with the model. However, there is only one replicate at 30 deg. C. More data is required to prove the hypothesis.

Table 16: The measured and calculated temperature dependence of the D value and 12 D sterilization time for polyester suture loops inoculated with bacillus subtilis (BS) var niger spores (ATCC 19659) according to AOAC protocol. The initial spore population was at least 1E6 CFU per carrier

Temp. deg. C	No. of Replicate Tests	Total No. of Carriers Tested	D value min. @ 60 mg/L	12CD hr. mg/L measured	12CD hr. mg/L <u>calculated</u> <u>from</u> <u>values @</u> <u>20 deg. C</u>	12D time minutes @ C = 60 mg/L
10	13	168	24	285	284	284
20	5	64	6	71	71	71
30	1	6	1.5	26	17.8	17.8
40					4.4	4.4
50					1.1	1.1

#### Materials Processing Method With Immersion Processing

##### Description - Fig. 7

Spin processing and spray processing configurations can generally provide better mass transport to the surface of the material to be processed than can immersion processing configurations. However, there are some applications where immersion processing can offer other advantages with respect to full wetting of complex surfaces. This can be particularly important in sterilization processing applications. In reference to Fig. 7, the outlet of three way valve 32 is connected by a short length of tubing to the process fluid inlet 48 of an immersion type materials processing module 50-I to a fill nozzle 36-F. The drain outlet 52B is connected by a short length of tubing to drain check valve 86. The outlet of check valve 86 is connected by a length of tubing to a drain line trap 54B. The drain line trap 54B is connected by a length of tubing to facility drain/reclaim 56. Purge gas supply 68 is connected by a short length of tubing to the inlet of two-way valve 70. The outlet of two-way valve 70 is connected to the inlet of sterile filter 82. The outlet of sterile filter 82 is connected to the module purge gas inlet 72. The overflow/waste gas vent outlet 52T is connected through a length of tubing to the inlet of a check valve 84. The outlet of the check valve 84 is connected

to the inlet of a trap 54T. The outlet of trap 54T is connected to the facility drain/reclaim for process effluent 56. The outlet of check valve 84 is also connected to the inlet of the catalytic ozone destruction unit 64. The outlet of the catalytic ozone destruction unit 64 is connected to a facility exhaust vent 66. The material(s) to be processed 38 may be in position inside the process module with a support structure 76.

#### Operation - Fig. 7

In reference to Fig. 7, the ozone gas-water solution is formed at a relatively low temperature and heated at the point-of-use as described earlier. Additional chemicals can be injected at the point-of-use from one or more chemical supplies as described earlier. The heated ozone-water solution flows from the outlet of three-way valve 32 through the process fluid inlet 48 in an immersion type material processing module 50-I to the a fill nozzle 36-F while the drain valve 86 is closed. Once the immersion module fills completely to the top with the heated ozone-water solution and the materials to be processed are immersed in the heated ozone-water solution, the heated ozone-water solution, together with any gases, flows out of the overflow /vent outlet 52T through filter 84, through the trap 54T to the facility drain/reclaim 56 for process effluent. Most of the undissolved gases that exit from the module pass through the catalytic ozone destruction unit 64 to the facility exhaust vent 66. The immersion processing module can be designed with baffles or other means to insure that the surfaces of the materials are continuously exposed to fresh heated ozone-gas-water solution as the solution is flowed from the module inlet 48 to the module overflow/vent outlet 52T. The module can be designed to hold instruments, devices, or materials with a minimum volume and thereby decrease the residence of the heated zone-water solution for a given ozone-water solution flow rate. The dissolved ozone concentration of the heated ozone-water solution will decay with time with a temperature dependent decay rate as discussed earlier. The residence time of the solution in the immersion module can be set at a predetermined value to insure that the dissolved concentration does not fall below a predetermined value for the duration of the cleaning or sterilization or oxidation process cycle. Once the cleaning cycle is complete the flow of the heated ozone-water solution to the process module may be stopped by setting the three-way valve 32 to direct the flow to the facility drain/reclaim 34 for the heated ozone-water solution. The module may be drained by opening the drain valve 78 and opening the two-way purge 70 to admit nitrogen, air, or some other purge gas into the module to replace the liquid that is drained from the module through check valve 86, through trap 54B to facility drain/reclaim 56. If the module is designed for sterile processing, then the filter 82 may be 0.2 micron filter to remove microbes from the purge gas. If the module is designed for ultra-



clean processing, then the filter 82 may be designed to remove particulate contamination from the purge gas. It is desirable to design the immersion processing module with baffles and other means to insure that the flow of liquid from the inlet to the outlet is approximately plug flow to insure that most of the "reduced concentration" or "spent" ozone-water solution and oxidation or cleaning byproducts are swept from the module and replaced by a "fresh" heated ozone-water solution. An alternative method of insuring that the "reduced concentration" or "spent" ozone zone-water solution and oxidation or cleaning byproducts are removed from the module and replaced by a "fresh" heated ozone-water solution is to fill process the materials with a series of fill, hold, drain, fill, hold, drain cycles. The module may be drained and refilled once the dissolved ozone concentration has fallen to a predetermined level. The solution may be stirred or mixed during the holding period with a high flow recirculation pump, with a stirrer, with ultrasonic or megasonic transducers, or other means known to those skilled in the art. Since the dissolved concentration of the heated ozone-water solution falls more quickly at higher temperatures, optimum hold times will typically be shorter at higher processing temperatures.

#### High Throughput Materials Processing Method With Multiple Single-Wafer Spin Processors Description - Fig. 8

The spin processing method can be implemented in a high throughput configurations for processing semiconductor wafers and the like. A single wafer processing tool can designed using an industry standard cluster tool geometry with multiple single-wafer process modules. Multiple single wafer spin processing modules may be served by a single ozonated water supply 22. In reference to Fig. 8, an ozonated water supply 22 is connected through a length of tubing to the common input of a three-way valve 24. The one outlet of three-way valve 24 is connected by a length of tubing to a manifold 23. Each of the six outlet ports of the manifold is connected by a length of tubing to each of six three-way valves 24. The three-way valves are labeled 24A, 24B, 24C, 24D, 24E, 24F. Three-way valve 24A is connected through a length of tubing to the cold process fluid inlet of a heat exchanger 28A. The other outlet of the three-way valve 24A is connected through a length of tubing to the facility drain-reclaim 26. The recirculating heating unit 30 which serves each of the heat exchangers is not shown. The heated process fluid outlet of heat exchanger 28A is connected through a short length of tubing to the inlet of a chemical injector/mixer 40A. The outlet of the injected chemical supply 42A is connected by a length of tubing to the inlet of a two-way valve 44A. The outlet of two-way valve 44A is connected by a length of tubing to the injection port of chemical injector/mixer 40A. The outlet of chemical injector/mixer 40A is

connected to the inlet port of three-way valve 32A. The one outlet of three-way valve 32A is connected through a short length of tubing to the etch inlet of a three-way valve etch/rinse valve 46A. The other outlet of the three-way valve 32A is connected by a length tubing to the facility drain/reclaim 34. The common outlet of three-way etch/rinse valve 46A is connected through a short length of tubing to the process fluid inlet 48A to a dispense nozzle (not shown) located in a gas tight materials processing module 50A. A rinse water supply 74 is connected by a length of tubing to a manifold 73. Each of the six outlet ports of manifold 73 is connected by a length of tubing to the rinse inlet of each of six three-way valves 46A, 46B, 46C, 46D, 46E, and 46F. Each of the other spin modules 50B, 50C, 50D, 50E, and 50F is connected in the same way. In the preferred embodiment each of the materials processing modules 50 may be fitted with a lid with a gas tight seal (not shown). The sealed materials processing module serves to contain the any ozone gas that is released from solution at the point of application. In the arrangement shown all modules share a single ozonated water supply and a single rinse water supply. Each spin processing module is shown with its own injected chemical supply. In an alternative embodiment, a single injected chemical supply may be used to supply injected chemicals to all modules.

The complete system may have a nitrogen purge supply 68, and each spin processing module will have a dispense nozzle 36, wafer chuck 58, spinner motor 60, drain outlet 52, vent outlet 62, purge gas inlet 72, purge gas two-way valve 70 as shown in Fig. 5 for a single module configuration. Each drain outlet 52 may be connected through a trap 54 to a facility drain 56 for process effluent as shown in Fig. 5 a single module configuration. Each vent outlet may be connected through a catalytic ozone destruction unit 64 to a facility waste gas exhaust vent 66 as shown in Fig. 5. These elements are omitted from Figure 8 for clarity. In one embodiment, a heat exchanger 28 may be used to increase the temperature of the ozone-water solution at the point-of-use. In another embodiment, a point-of-use water heater 29 may be used in lieu of a heat exchanger.

### High Throughput Materials Processing Method With Multiple Single-Wafer Spin Processors - Multiple Heat Exchangers

#### Operation - Fig. 8

The operation of each wafer spin processing module and associated elements is the same as described earlier under Fig. 5. The modules can be operated by a process controller (not shown) to achieve high throughput while sharing the most expensive resource, the ozonated water supply. A typical process sequence for such a tool is shown in Table 13. A

single ozone-water supply feeds the three process modules that are in the etch/clean phase at any point in the process cycle. A process controller controls the flow of ozone-water solution, injected chemicals, and DI rinse water to each of the modules by controlling the positions of the control valves. In all the embodiments shown in all the figures the valves may be gas operated or electrically operated solenoid valves so that the valves can be controlled with a central controller. In this example the etch time is set to 60 seconds and the rinse, dry, and load/unload times are each set to 20 seconds. The throughput for the example shown is 180 wafers/hour. During any 20 second period, 3 modules at most are in an etch/clean cycle, 1 module at most is in the DI water rinse cycle, and 1 module at most is in the Spin Dry Cycle. If for example the ozonated water supply flow rate required for each module during the etch/clean cycle is 3.33 liters per minute at a given dissolved ozone concentration, then the entire cluster of 6 modules may be supplied by a single ozonated water supply with a flow capacity of 10 liters per minute. The example operating sequence in Table 17 shows the operation during steady state when three process modules are etching, one process module is rinsing, one process module is drying, and one process module is being unloaded, and loaded with the next wafer to be processed. During the period when each of the modules of the cluster are loaded in succession for the first time at the beginning of a run of wafer, first only one processor is in the etch mode, then two processors are in the etch mode, then three processors are in the etch mode. There is an analogous transient period when each of the modules of the cluster are unloaded in succession at the end of a run of wafers. At the end of a run of wafers, three processors are in the etch mode, then two processors are in the etch mode, then one processor is in the etch mode. During the period in which only one module or only two modules are in the etch mode, only 3.33 L/min or only 6.66L/min of ozonated water is being used for etching for this example. In one embodiment the total ozonated flow can be held constant (~10.0 L/min for this example) either by setting the three-way valves 24, or by setting the three-way valves 32 to direct to the facility waste/reclaim 34 the ozonated water not needed for etching. This approach maintains a steady state water flow through the ozone-gas-water contactor and provides a simple method to maintaining a stable dissolved ozone concentration. This approach is simple. An alternative method is to reduce the flow rate of ozonated water supplied by the ozonated water supply 22 while maintaining the same supplied dissolved ozone concentration. This requires that the ozonated water supply include a controller for maintaining the same dissolved ozone concentration at different flow rates less than the maximum flow rate for that concentration. This approach is more challenging because it would require that the controller be able to quickly stabilize the concentration in

about 2 seconds or less, as the flow rate was incremented in steps at the start of the process run, and decremented in steps at the end of a process run.

Table 17: Operation of Cluster Tool with Six Process Modules - Example

Module	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec	20 sec
#A	L/U	E	E	E	R	D	L/U	E	E	E	R	D	L/U	E	E	E	R	D	
#B	D	L/U	E	E	E	R	D	L/U	E	E	E	R	D	L/U	E	E	E	R	
#C	R	D	L/U	E	E	E	R	D	L/U	E	E	E	R	D	L/U	E	E	E	
#D	E	R	D	L/U	E	E	E	R	D	L/U	E	E	E	R	D	L/U	E	E	
#E	E	E	R	D	L/U	E	E	E	R	D	L/U	E	E	E	R	D	L/U	E	
#F	E	E	E	R	D	L/U	E	E	E	R	D	L/U	E	E	E	R	D	L/U	
E	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	
R	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
D	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
L/U	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	

E - Etch/Clean Cycle

R- DI Water Rinse Cycle

D- Spin Dry Cycle

L/U - Load/Unload Wafer w/ Robot

Cluster Throughput for Example: 30 wafers/hour/spinner x 6 spinners=180 wafers/hour

### High Throughput Materials Processing Method With Multiple Single-Wafer Spin Processors - Single Shared Heat Exchanger

#### Description - Fig. 9

In reference to Fig 9, an alternative embodiment may use a single point-of-use heater (heater exchanger 28 or point-of-use heater 29) and single chemical injector/mixer 40 to serve

multiple spin processing modules. Since the flow rate of the ozone-water solution through the point-of-use heater is higher, the heat input required to increase the temperature from T1 to T2 is higher. The heat exchanger may be larger and the volume of the heat exchanger may be larger. The residence time in the heater and connecting tubing may be kept approximately the same if the volume is increased by the same factor as the flow rate. In the example shown in Table 17 above, the flow rate through the exchanger and injector/mixer will be up to 3 times higher and the power input required will be up to 3 times higher.

#### Operation - Fig. 9

The operation of the embodiment shown in Fig. 9 is otherwise similar to that described under Fig. 8 and Fig. 5. In the embodiment using the single heat exchanger, the flow of heated ozone-water solution is directed to the three spin processing modules that are in the etch phase of the cycle. During the period when each of the modules of the cluster either are loaded in succession for the first time at the beginning of a run of wafers, or are loaded in succession for the last time at the end of a run of wafers, fewer than three modules may be in the etch mode at one time as described under Fig. 8. In one embodiment the total ozonated flow can be held constant (~10.0 L/min for this example) by setting the three-way valves 32 to direct to the facility waste/reclaim 34 the ozonated water not needed for etching. This approach maintains an approximately steady-state water flow through the ozone-gas-water contactor and provides a simple method to maintaining a stable dissolved ozone concentration as described under Fig. 8. It also maintains a constant flow rate through the shared heat exchanger 28 and maintains a constant dispense temperature. This is again a simple embodiment. An alternative method is to reduce the flow rate of ozonated water supplied by the ozonated water supply 22 while maintaining the same supplied dissolved ozone concentration and same dispense temperature. This embodiment requires that the ozonated water supply include a controller for maintaining the same dissolved ozone concentration at different flow rates as discussed under Fig. 8. This also requires the use of a point-of-use heater 29 with the means to maintain a constant dispense temperature at different dispense flow rates.

#### High throughput materials processing method with an on-axis wafer-batch spin processor Description - Fig. 10

The spin processing method can be implemented in a high throughput configuration for processing semiconductor wafers and the like. A batch wafer processing tool can be designed using an on-axis batch spin processor designed to process two to four cassettes of wafers or

substrates at one time. In reference to Fig. 10, a batch spin module 50-B1 for processing two cassettes of wafers in an on-axis spin configuration may be served by a source of ozonated water, rinse water, purge gas, drain, and vent connections using the same configuration as that described under Fig. 5 for a single wafer spin processor. In the case of the on-axis wafer-batch spin processor, the outlet of the three-way etch rinse valve 46 is connected by a short length of tubing through a module inlet 48 to a perimeter spray bar 36-PB. Two 25 wafer cassettes 90 and 92 are mounted in a wafer cassette chuck 94. The wafer cassette chuck 94 is connected by a shaft or other means to a spin motor 60. The spin motor may be set to a particular acceleration rate, spin RPM, spin duration, and deceleration rate under microprocessor control.

#### Operation - Fig. 10

In reference to Fig. 10, the operation of an on-axis batch spin processor designed to process two to four cassettes of wafers or substrates at one time is similar in most respects to the operation of a single wafer spin processor described under Fig. 5. A batch spin processor typically operates at a lower RPM and lower flow rate per wafer than a single wafer spin processor. The RPM for a batch spinner is typically in the range of 500 to 1500 RPM. The total etch chemistry flow rate for a batch wafer spinner is typically in the range of 10 to 20 liters/minute. The lower RPM and lower flow rates will yield a lower mass transport rate and lower etch rate. The temperature at which the etch rate will become mass transport limited will be lower since the mass transport rate is lower as discussed earlier.

In the etch mode a heated ozone water solution flows from the outlet of three-way valve through an inlet 48 in the module, to the perimeter spray bar 36-PB. The heated ozone-water solution is dispensed from multiple positions along the length of the spray bar and contacts the surfaces of the wafers held in the wafer cassettes which are spinning on-axis.

#### High throughput materials processing method with an off-axis wafer-batch spin processor Description - Fig. 11

The spin processing method can be implemented in a high throughput configuration for processing semiconductor wafers and the like. A batch wafer processing tool can be designed using an off-axis batch spin processor designed to process two to four cassettes of wafers or substrates at one time. In reference to Fig. 11, a batch spin module 50-B2 for processing two cassettes of wafers in an off-axis spin configuration may be served by a source of ozonated water, rinse water, purge gas, drain, and vent connections using the same configuration as that

described under Fig. 5 for a single wafer spin processor. In the case of the off-axis wafer-batch spin processor, the outlet of the three-way etch rinse valve 46 is connected by a short length of tubing through a module inlet 48 to a center spray tower 36-CT. Two 25 wafer cassettes 90 and 92 are mounted in a wafer cassette chuck 96. The wafer cassette chuck 96 is connected by a shaft or other means to a spin motor 60. The spin motor may be set to a particular acceleration rate, spin RPM, spin duration, and deceleration rate under microprocessor control.

#### Operation - Fig. 11

In reference to Fig. 11, the operation of an off-axis batch spin processor designed to process two to four cassettes of wafers or substrates at one time is similar in most respects to the operation of a single wafer spin processor described under Fig. 5. A batch spin processor typically operates at a lower RPM and lower flow rate per wafer than a single wafer spin processor. The RPM for a batch spinner is typically in the range of 500 to 1500 RPM. The total etch chemistry flow rate for a batch wafer spinner is typically in the range of 10 to 20 liters/minute. The lower RPM and lower flow rates will yield a lower mass transport rate and lower etch rate. The temperature at which the etch rate will become mass transport limited will be lower since the mass transport rate is lower as discussed earlier.

In the etch mode a heated ozone water solution flows from the outlet of three-way valve through an inlet 48 in the module, to the center spray tower 36-CT. The heated ozone-water solution is dispensed from multiple positions along the length of the spray tower and contacts the surfaces of the wafers held in the wafer cassettes which are spinning off-axis. The ozone-water solution that moves across the wafer, entering at an edge near the spray tower and exiting at an edge far from the spray tower. The dissolved ozone concentration falls as the solution moves across the wafer surface because the ozone in solution is depleted by reaction with the material on the wafer surface. In a standard off-axis batch processor configuration this can yields a relatively high etch rate at the wafer leading edge and a relatively low etch rate at the wafer trailing edge. A preferred embodiment for off-axis wafer processing not only rotates the wafer cassettes around an axis some distance form the wafer center, but also slowly rotates the wafer cassettes on axis so that all edges of the wafers are placed in the leading edge position for approximately the same time during the wafer processing cycle. This can be accomplished by a drive gear or other device for slowly rotating the wafer cassettes about the center of the wafers as the wafer cassettes are rotated off axis.

## 2<sup>ND</sup> GROUP: Heat Applied At The Point Of Application (Fig. 12 through Fig. 15)

A second method for oxidizing materials at high speed using a solution of ozone gas dissolved in water comprising the steps of dissolving (relatively high concentration) ozone gas in water at a relatively low predetermined temperature T1 to form an ozone-water solution (with a relatively high dissolved ozone concentration), applying the cold ozone-water solution to said materials while heating said materials and said ozone-water solution at the point of application to quickly increase said material temperature and said solution temperature to a predetermined higher temperature T2>T1 where preferably T2-T1> 5 degree C. The second method for oxidizing materials at high speed may additionally include a means for injecting and mixing additional chemicals into the ozone-water solution just upstream of the point of application of the ozone-water solution to the materials to be oxidized. A number of preferred embodiments are illustrated in Fig. 12 to Fig. 16.

### Material to be Processed and Ozone-Water Solution Heated at the Point of Application - Top Surface Radiant Heating

#### Description - Fig. 12

This embodiment is very similar to the embodiment shown in Fig. 5 except that the ozone water solution is not heated upstream of the point-of-use. Instead, the material to be processed and the ozone-water solution is heated at the point of application. Accordingly, the point-of-use water heater is omitted and point of application heater is added. With reference to Fig 12, an ozonated water supply 22 is connected through a length of tubing to the common input of a three-way valve 24. The one outlet of three-way valve 24 is connected through a length of tubing to the process fluid inlet of chemical injector/mixer 40. The other outlet of three-way valve 24 is connected through a length of tubing to the facility drain-reclaim 26 for the ozone-water solution. The outlet of an injected chemical supply 42 is connected to the inlet of two-way valve 44. The outlet of valve 44 is connected to the chemical injection port of chemical injector/mixer 40. The outlet of chemical injector/mixer 40 is connected by a short length of tubing to the common inlet port of three-way valve 32. The one outlet of three-way valve 32 is connected through a short length to one inlet of a three-way valve etch/rinse valve 46. The common outlet of three-way etch/rinse valve 46 is connected through a short length of tubing to the process fluid inlet 48 to a dispense nozzle 36 located in a gas tight materials processing module 50. The materials processing module 50 is fitted with a lid with a gas tight seal with a quartz window 98 in the lid with good infrared



transmission. The module purge gas connection 72, the module drain connection 52, and module vent connections 62 are the same as described previously. A high power radiant heat source 100 such as a quartz halogen lamp array is positioned to illuminate the surface of the wafer or substrate 38 which is mounted on a spinner chuck 104. We can calculate an estimate of the maximum amount of power required to heat the material and the ozone water solution by assuming that we must increase the temperature of the full flow of ozone water solution as it passes over the wafer and we must heat the full volume of the wafer. The power required increase the water temperature by a given amount for a given flow rate has been presented in Table 5 of the discussion under Fig. 1. The power required to increase the temperature of an ozone water solution flowing at 2.7 L/min by 50 degree C is approximately 10 Kwatts. A small amount of additional energy will be required to increase the wafer temperature by the same amount. The amount of energy can be readily calculated from the mass and heat capacity of the wafer or substrate. If in practice the only the surface of the wafer or only a thin layer of the ozone-water solution at the wafer-solution interface is heated, then the power requirements will be lower. Also, if the ozone-water solution flow rate is reduced, the power requirement will be lower. The methods of heating of wafers and substrates with quartz halogen lamp arrays are well developed and well known to those skilled in the art in the area of single wafer rapid thermal processing (RTP). A typical RTP lamp array for 200 mm diameter wafers has a power input of about 50 Kwatts. Typically about 40 percent of the lamp input power (20 Kwatts) is deposited in the substrate in RTP applications. The RTP lamp power is well suited to this new application. A gold coated infrared reflective surface 102 is place above the radiant heat source 100 to reflect the visible and infrared radiation back to the substrate. A gold coated infrared reflective surface is placed on the surface of the spinner chuck to to reflect the visible and infrared radiation that may pass through the substrate back though the substrate toward the infrared reflective surface 102. The length of tubing leading to the dispense nozzle 36 may be insulated and protected with an infrared reflective surface like gold to prevent the tubing and dispense nozzle from overheating. A non-contact temperature sensor 106 is positioned to read the temperature of the substrate. The measurement of the temperature of spinning silicon wafers is well known to those skilled in the art in the area of single wafer rapid thermal processing (RTP).

### Operation - Fig. 12

A cold ozone-water solution at temperature  $T_1$  is applied to a material to be processed while the material to be processed and the ozone-water solution is heated to temperature  $T_2 > T_1$  at the point of application using radiant heating from the top surface. In one embodiment the ozone water solution is dispensed onto the wafer surface with a center dispense nozzle as discussed earlier under Figure 5. The cold ozone-water solution that flows radially outward to the edge of the spinning wafer will cool the wafer when the radiant heat source is off. Temperature sensor 106 is connected to controller (not shown) which adjusts the heat output of the radiant heat source 100 to maintain a preset temperature (50 degree C for example). The use of feedback control to control of the temperature of wafers and other substrates heated with quartz halogen radiant heating sources is also well known to those skilled in the art in the area of single wafer rapid thermal processing (RTP). When the radiant heat source is turned on to quickly increase the temperature of the substrate the substrate will be heated by absorption of radiant energy. The ozone-water solution that passes across the surface of the wafer or substrate in a thin layer will not only absorb some radiant energy, but will be heated by convection as the water passes over the heated substrate and energy is transported from the heated substrate to the chilled ozone-water solution. Since both velocity and thickness of the water that moves radially outward varies with radius, both the rate of energy loss by water cooling and the rate of energy gain by radiant heating and absorption may have a radial dependence. Accordingly, the radiant heat source may be designed to deposit a different amount of energy into different radial zones to achieve approximately the same temperature over the entire substrate surface. The use of concentric zone radiant heating with multiple temperature sensors to maintain a constant temperature over the entire substrate surface is again well known to those skilled in the art in the area of single wafer rapid thermal processing (RTP).

The ozone-water solution will not be heated until the ozone-water solution is applied to the surface of the substrate. Accordingly, the time delay between the increase in temperature of the ozone-water solution and the application of the solution to the surface of the material to be processed will be essentially zero, the rate at which the ozone gas leaves the solution at the substrate should not differ significantly from the case in which the substrate is not heated but is at the same temperature as the ozone-water solution. This type of embodiment will be particularly helpful in preserving a high dissolved ozone concentration in embodiments in which the optimum processing temperature is higher because the concentration decays in shorter time at higher temperatures as discussed earlier. A solution

application means which achieves a higher mass transport coefficient  $M$  has a higher maximum optimum processing temperature.

#### Material to be Processed and Ozone-Water Solution Heated at the Point of Application - Bottom Surface Radiant Heating

##### Description - Fig. 13

In reference to Fig. 13, the structure of this embodiment is similar to that described under 12 except that the wafer or substrate is mounted on an edge-driven open-bottom spinner-chuck 108, and the wafer is heated from the bottom. A radiant heat source 100 illuminates the bottom of the wafer through a quartz window 98 in the bottom of the spinner module. An infrared reflective surface 102 is placed behind the radiant heat source and a second infrared reflective surface 102 is placed above the dispense nozzle 36.

##### Operation - Fig. 13

In reference to Fig. 13, the operation of this embodiment is similar to that described under 12 except that the wafer or substrate is heated with a radiant heat source 100 which illuminates the bottom of the wafer. This configuration may be more convenient to implement in some applications.

#### Material to be Processed and Ozone-Water Solution Heated at the Point of Application - Bottom Surface Convection Heating

##### Description - Fig. 14

In reference to Fig. 14, the structure of this embodiment is similar to that described under 13 except that wafer is heated from the bottom convectively. The substrate is mounted on an edge-driven open-bottom spinner-chuck 108. A source of heated fluid 112 is dispensed from a nozzle 114 to transfer heat to the bottom of the wafer. The heated fluid may be a liquid such as hot water or a hot gas such as nitrogen. The heated fluid may be produced by passing the fluid through a point-of-use heater.

##### Operation - Fig. 14

In reference to Fig. 14, a cold ozone-water solution at temperature  $T_1$  is applied to a wafer or substrate 38 to be processed with nozzle 114 while the material to be processed and the ozone-water solution is heated to temperature  $T_2 > T_1$  at the point of application through the use of a heated fluid at a temperature greater than  $T_2$  directed at the bottom of the substrate with a nozzle 114. The rate at which the substrate can be heated is determined by the net rate

of energy gain. The net energy gain is the energy that is transferred to the wafer from the heated fluid stream incident upon the bottom of the substrate minus the energy lost from the wafer to the cooled ozone water stream incident upon the top of the substrate.

#### Material to be Processed and Ozone-Water Solution Heated at the Point of Application - Bottom Surface Conduction and Convection Heating

##### Description - Fig. 15

The structure is similar to that described under Fig.14 except the wafer is mounted on a heated spinning wafer chuck 116 which is driven by spinner motor 60. In reference to Fig. 15, a wafer or substrate 38 is mounted on a heated spinning wafer chuck 116. A non-contact temperature sensor is positioned to read the temperature of the top surface of the substrate. The temperature sensor 106 is connected to a controller. The controller adjusts the amount of energy transferred to the wafer by the heated chuck by adjusting the temperature of the chuck surface.

##### Operation - Fig. 15

A cold ozone-water solution at temperature  $T_1$  is applied to a material to be processed while the material to be processed and the ozone-water solution is heated to temperature  $T_2 > T_1$  at the point of application using a heated chuck in contact with or close proximity to the bottom side of the wafer to heat the wafer.

The embodiments shown in Fig. 12 and Fig. 13 are usually preferred over those shown in Fig. 14 and Fig. 15 because they are based upon a well developed technology and because they can transfer the most power to the wafer or substrate and thereby achieve very fast heating rates. The embodiment shown in Fig. 15 may be lowest cost. However, the embodiment in Fig. 15 may not be able to transfer sufficient energy to the substrate to overcome the energy loss to the chilled ozone-water solution. This heating embodiment may be capable of providing sufficient energy transfer when used in an application for which the flow rate of the chilled ozone-water solution to the substrate is very low.

#### Material to be Processed and Ozone-Water Mist Solution Heated at the Point of Application - Top Surface Radiant Heating

This embodiment differs from all the embodiments presented in Fig. 1 to Fig. 15 in an important respect. In this embodiment a chilled ozone-water mist solution mixed with ozone gas is applied to the surface of a spinning wafer or substrate. This embodiment has the

potential to achieve higher removal rates than prior art moist ozone gas methods. The dissolved ozone concentration in the chilled ozone-water mist has the potential to be higher than that which can be produced at the higher temperatures, 50 to 95 degree C, employed in prior art moist ozone gas methods. This embodiment may be able to achieve relatively high removal rates at the low water flow rates usually associated with prior art moist ozone gas organic removal methods.

#### Description - Fig. 16

In reference to Fig. 16, a ozonated water mist/ozone gas supply 118 (detail described in Fig. 19) connected by a short large diameter (approximately 20 cm long, 10 cm diameter) length of tubing to a gate valve 122 to a process fluid inlet 126 in the materials processing module 50-OM. The ozone gas/ozonated water mist outlet 62 of the materials processing module is connected by a short length (~10cm long, ~10 cm diameter) of tubing to a mist eliminator 124. The outlet of mist eliminator 124 is connected to ozone catalytic destruction unit 64. The outlet of the ozone catalytic destruction unit 64 is connected to facility exhaust vent inlet 66. The liquid drain outlet 52 of the process module 50-OM is connected by a length of tubing through a trap 59 to the facility drain/reclaim inlet 56 for process effluent. A purge gas supply 68 is connected by length of tubing through a two way valve 70 to a purge gas inlet 72 in the process module. A wafer or substrate is mounted on a wafer chuck 104. The wafer chuck is attached to a spinner motor 60. The wafer chuck 104 has an infrared reflective surface to prevent heating of the chuck by the radiant heat source. The process module 50-OM is fitted with a quartz window 98. A radiant heat source 100 and Infrared reflective surface 102 is positioned to illuminate the wafer surface as described under Fig. 12. A non-contact temperature sensor 106 is positioned to read the temperature of the wafer as described under Fig. 12. A rinse water supply 74 is connected by a length of tubing through a two-way valve 120, through the rinse water inlet 128 in the process module to a rinse water dispense nozzle 36 positioned to dispense rinse water on the wafer surface.

### Operation - Fig. 16

In reference to Fig. 16, a cold ozone-water mist solution at temperature T1 mixed with ozone gas, is applied to a material to be processed while the material to be processed and the ozone-water mist solution is heated to temperature  $T2 > T1$  at the point of application using radiant heating from top surface. The radiant heating system has been described under Fig. 12. At the end of the spin etch clean cycle, the gate valve 122 is closed and the rinse valve 120 is opened to spin rinse the wafer. At the end of the spin rinse cycle, the rinse valve is closed and the wafer is spun dry.

### 3<sup>rd</sup> GROUP: High Performance Ozonated Water Supply for Use in Conjunction with the Preferred Embodiments (Fig. 17 through Fig. 19)

A method of providing a supply of an ozone-water solution with a relatively high dissolved ozone concentration comprising the steps of chilling water to a relatively low predetermined temperature, flowing the chilled water and high concentration ozone gas through into an ozone gas-water contactor to form an ozone-water solution, flowing the undissolved ozone gas out of the contactor, flowing the ozone-water solution out of the contactor to form a supply of ozonated water. The contactor is designed to provide a relatively large surface area to provide for a high rate of mass transfer of the ozone gas into water solution. A relatively large contacting volume can provide sufficient time for a significant portion of the ozone gas to be transported into solution. Preferred embodiments are illustrated in Fig. 17 and Fig. 18. By reducing the DI water temperature from 20 degree C to 5 degree C the saturation concentration is increased by a factor of two.

A method of providing a supply of an ozone-water-mist solution with a relatively high dissolved ozone concentration comprising the steps of chilling water to a relatively low predetermined temperature, flowing the chilled water mist through an liquid atomizer to form a chilled water mist, flowing the chilled water mist and high concentration ozone gas through a contacting volume to supply an ozone-water solution mixed with ozone gas at the outlet of the contactor volume. The chilled water mist with very small droplet size provides a relatively large surface area to provide for a high rate of mass transfer of the ozone gas into water solution. The a relatively large contacting volume can provide sufficient time for a significant portion of the ozone gas to be transported into solution. A preferred embodiment is illustrated in Fig. 19.

## Typical Ozone-Water Solution Supply Using Venturi Injector And Downstream Bubble Column

### Description - Fig. 17

In reference to Fig. 17, a DI water supply 130 is connected through a length of tubing to the fill inlet of a three-way valve 132. The outlet of three-way valve 132 is connected through a length of tubing to the inlet of pump 134. The outlet of pump 134 is connected to the process fluid inlet of heat exchanger 136. A recirculating cooling unit is connected to the working fluid inlet and outlet of the heat exchanger 136. The process fluid outlet of the heat exchanger 136 is connected to the motive flow inlet of venturi injector 140. The outlet of venturi injector 140 is connected to a bubble column inlet 142 in bubble column 144. The pump is a positive displacement pump capable of delivering the pressure to overcome the pressure drop across the exchanger 136 and venturi injector 140. The supply outlet 156 of the bubble contactor 146 is connected by a length of tubing to a liquid back pressure regulator 158. The outlet of liquid back pressure regulator 158 is connected by a length of tubing to two-way valve 160. The outlet of the two-way valve 160 is connected to the process to provide a pressurized source of ozonated water at 162.

An pressurized oxygen supply 164 is connected by a length of tubing to a two-way valve 166. The outlet of two-way valve 166 is connected by a length of tubing to mass flow controller 168. The outlet of mass flow controller 168 is connected to the oxygen gas inlet of high concentration ozone generator 170. A recirculating cooling unit 172 is connected to cool the ozone generator cells of ozone generator 170. The ozone gas outlet of ozone generator 170 is connected to the inlet of check valve 174. The outlet of check valve 174 is connected to the suction gas inlet of venturi injector 140. The waste ozone gas vent outlet 176 of contactor 146 is connected to an ozone catalytic destruction unit 178. The outlet of ozone catalytic destruction unit 178 is connected to the inlet of off-gas back-pressure regulator 180. The outlet of back pressure regulator 180 is connected to the inlet to the facility vent 182 for contactor off-gas. The recirculate-outlet 154 of contactor 146 is connected by a length of tubing to the recirculate inlet of three-way valve 132. An level sensor 148 is positioned several inches from the top of the contactor 146 to detect the position of the minimum liquid level in the contactor. An level sensor 150 is positioned a short distance (1- 2 inches) above level sensor 148 to detect the position of the maximum liquid level in the contactor. The level sensors are connected to a fill level controller 152. The fill level controller connected to the fill valve 132, a solenoid valve, to control the filling of the contactor as water is removed from the contactor at the supply point 162. The connection from the controller to the valve is not

shown. This embodiment is designed to provide for dissolving ozone gas in water at a pressure above atmospheric to increase the dissolved ozone concentration as predicted by Henry's law. Since the contactor and connecting tubing is chilled below ambient, the entire system is insulated.

In an alternative embodiment, designed to dissolve ozone gas at atmospheric pressure, the liquid back pressure regulator 158 may be eliminated and the gas back pressure regulator 178 is moved to outlet line of the ozone generator 170 to maintain the required pressure in the ozone generator for efficient generation of ozone gas. In this alternative embodiment the contactor is not pressurized. Accordingly, a pump (not shown) must be placed between the contactor supply outlet 156 and the valve 160 to provide a pressurized source of ozonated water at 162. In this case the two-way valve solenoid valve (all valves are understood to be solenoid valves) 160 may be replaced by a three way solenoid valve (not shown) with the normally open outlet connect by a length of tubing back to the contactor 146 and the normally closed outlet connected to 162. In this configuration, the ozonated water can be recirculated back to the contactor (connection not shown) to keep the concentration in the line a full concentration, and then dispensed to the process by actuating the three-way dispense valve (not shown). In electronic device cleaning/processing applications, all wetted materials may be Teflon PFA or Teflon PTFE materials in lieu of metal materials to exclude the introduction of metal contamination into the process.

#### Operation - Fig. 17

In reference to Fig. 17, DI water at near room temperature flows through the heat exchanger 134 where it may be cooled to a lower temperature (5 degree C for example). The chilled DI water enters the venturi injector 140 where ozone gas is injected. Oxygen gas under pressure is regulated to a delivery pressure (40 psig. for example) and supplied to the inlet of mass flow controller 168. The mass flow controller controls the mass flow to the inlet of the ozone generator 170. An ozone generator such as the ASTEX AX8100 can deliver ozone at a concentration of about 240 g/NM<sup>3</sup> at a flow rate of 1.5 L/min. An ASTEX AX 8200 can deliver ozone at a concentration of about 240 g/NM<sup>3</sup> at a flow rate of 4.5 L/min. The ozone gas enters the suction inlet of venturi injector 140 where the high shear forces produce many very small bubbles of ozone gas which move to the bubble contactor with the water flow. The chilled DI water, mixed with ozone gas bubbles, moves through the contactor as shown by the arrows on Fig. 17. The large surface area of the many small bubbles provide for a high rate of mass transfer from the gas to the liquid by diffusion. The residence time of



the bubbles in the contactor determines the concentration that can be achieved at the end of one pass through the contactor. The ozone gas that does not dissolve into the water exits from the contactor at the outlet 176. The level controller 152 controls the opening and closing of the fill valve 132. The level of water in the contactor is controlled as water is drawn from the contactor at 162.

An ozonated water supply can be designed to operate either as a batch recirculating dissolved ozone supply system or as a single pass continuous flow dissolved ozone supply system. Both systems have a means for generating high concentration ozone gas, a means for cooling a volume of water, and a means for dissolving the ozone gas in the water. The key design elements are summarized in Table 18.

Table 18. Key Design Elements: Batch Recirculating Compared to a Continuous Flow Ozonated Supply

Design Element	Batch	Continuous
	Ozone Mass Flow Out > Ozone Mass Flow Dissolved	Ozone Mass Flow Out = Ozone Mass Flow Dissolved
supply duty cycle	intermittent	continuous
ozone generator flow	variable	constant
req'd ozone gas mass flow rate	< DIO3 mass flow rate	≥ DIO3 mass flow rate
operating temperature	< 20 C or ≥ 20 C	< 20 C or ≥ 20 C
req'd DI water cooling capacity	proportional to average DIO Flow	proportional to peak DIO Flow
operating pressure	P=1 bar or P> 1 bar	P=1 bar or P> 1 bar
gas-liquid contactor element	venturi injector or packed column	venturi injector or packed column
gas-liquid contact volume	bubble column or packed column	bubble column or packed column
ozone injection pressure	<1bar or contactor pressure	<1bar or contactor pressure
maximum DIO3 concentration	Cout = saturation concentration	Cout < saturation concentration
system cost	lowest	higher

Let us describe the operation of small batch type ozonated supply system of the type employed in the inventors laboratory. A small venturi injector and bubble column contacting system for dissolving ozone gas at one atmospheric pressure employ a high pressure pump 134 feeding a small venturi injector 140 (Mazzei Injector Model 287) with chilled DI water at a motive flow rate of 3.5 L/min. (See Fig. 17) The water enters the inlet 142 in the center of the internal bubble column 144, then spills over the partition to the outer annular volume between the column 144 and the internal wall of the contactor 146 and returns to the pump by exiting from the outlet 154. Accordingly, the pump 134 continuously circulates water in a closed loop from the contactor 146, through the venturi injector 140 at a flow rate of about 3.5 L/min, and back to contactor bubble column 144.. Once refilled with fresh water, the concentration can be restored to saturation in about 20 to 30 minutes for a ozone flow rate of 0.48 L/min at a gas phase ozone concentration of 240 g/NM3 and a contactor volume of 12 liters.

A pressure of about 40 psig is required to pass a motive flow of 3.5 L/min through the venturi in a system with an unpressurized contactor ( $p=1\text{bar}=14.5\text{psia}$ ). Ozone gas from the ozone gas delivery system is fed to the suction inlet port of the venturi at a flow rate of 0.5 Liters/min. The pressure at the suction port of the venturi is about 0.7 to 0.8 bar (aprox. 10 to 12 psia). The injection conditions are summarized in Table 19 below. In a recirculating system of this type, the dissolved ozone concentration in the contactor starts from zero and rises exponentially to a maximum saturation value which is determined by Henry's law given the ozone gas concentration and pressure and the water temperature. In an unpressurized system, the ozone gas is dissolved at 1 bar=14.5 pisa. For the flow rates and concentrations specified, the exponential time constant with which the dissolved ozone concentration approaches saturation is about 6 to 7 minutes. Accordingly, one can produce a 12 liter batch of ozonated water at a concentration of about 95% of saturation is 20 minutes (about three time constants). The key design parameters for this small capacity batch system are summarized in Table 19. Much larger capacity systems with higher mass flow rates can be designed using the same approach.

Table 19 Flows, Pressures, and Venturi Injector Size for a venturi-based ozone gas-water contacting system for dissolving ozone gas at a pressure 1 bar and a flow rate of 0.48 L/min. The predicted saturation concentration with a water temperature of 5 degree C and ozone gas concentration of 240 g/Nm<sup>3</sup> is also shown

PARAMETER	Injector Design Values
venturi outlet/contactor pressure (bar)	1 bar
venturi outlet pressure – pisa (psig)	43.5 (28.8)
Injector (ozone) flow - liters/min (ft <sup>3</sup> /hr)	.48 (1.0)
Venturi Inlet Pressure - psig	40
Motive (water) flow - liters/min (gpm)	3.5 (.8)
Pump Pressure Boost Required - psig	40
Venturi Size (Mazzei Model Number)	Model 287
Predicted Saturation Concentration	109 mg/L

If the mass flow rate of the ozone to the injector is increased by increasing the ozone gas flow rate at a given gas phase concentration, then the volume of water will approach saturation in a shorter time. For example, the Astex AX 8100 ozone generator can supply ozone at a concentration of about 240 g/Nm<sup>3</sup> at a flow rate of 1.5 L/min at a generator power setting of 90%. An Astex AX8200 ozone generator has three times the capacity of an AX8100 and can supply ozone at a concentration of about 240 g/Nm<sup>3</sup> at a flow rate of 4.5 L/min at a generator power setting of 90%. The mass transfer rate from the gas to the liquid is a function of the surface area and the concentration difference between the gas phase and liquid phase. Accordingly, the mass transfer rate gradually falls as the dissolved concentration rises.

The entire ozone gas-water contacting system may be enclosed in a 19 inch equipment rack equipped with a ventilation system and a safety interlock system employing ozone leak detection sensors. In a system designed for use in semiconductor applications all the wetted parts can be Teflon or PVDF to eliminate iron contamination associated with stainless steel components.

The contactor can be either pressurized or unpressurized. If the contactor is designed to withstand pressures above one atmosphere, then the ozone gas can be dissolved in water at elevated pressures and thereby produce a higher dissolved ozone concentration at a given ozone gas phase concentration and water temperature than can be achieved at one atmosphere.

For example, ozone can be dissolved in water at 4 bar (58 psia) In this case the pump provides boost pressure above the contactor pressure and the venturi is selected to achieve adequate suction at desired ozone flow rate. In Table 20 below we have presented the parameters for a high pressure venturi-based ozone gas-water contacting system for dissolving ozone gas at a pressure 3, or 4 bar and a flow rate of 1.5 L/min or 4.5 L/min. In such a system the off gas from the contactor must exit from the pressurized contactor through a back pressure regulator set to the desired contactor pressure and the liquid stream that is dispensed to the process must also exit from the contactor through a back pressure regulator set to the desired contactor pressure as shown in Fig. 17.

Table 20 Flows, Pressures, and Venturi Injector Size for a venturi-based ozone gas-water contacting system for dissolving ozone gas at a pressure 3, or 4 bar and a flow rate of 1.5 L/min or 4.5 L/min. The predicted saturation concentration with a water temperature of 5 degree C and ozone gas concentration of 240 g/Nm<sup>3</sup> is also shown

PARAMETER	Injector Design Values			
	3 bar	4 bar	3 bar	4 bar
venturi outlet/contactor pressure (bar)	3 bar	4 bar	3 bar	4 bar
venturi outlet pressure – pisa (psig)	43.5 (28.8)	58 (43.3)	43.5 (28.8)	58 (43.3)
Injector (ozone) flow - liters/min (ft <sup>3</sup> /hr)	1.5 (3.2)	1.5 (3.2)	4.5 (9.6)	4.5 (9.6)
Venturi Inlet Pressure - psig	70	100	90	120
Motive (water) flow - liters/min (gpm)	14.5 (3.8)	17.2 (4.5)	29.5 (7.8)	35.6 (9.4)
Pump Pressure Boost Required - psig	41.2	56.7	46.5	76.7
Venturi Size (Mazzei Model Number)	Model 484	Model 484	Model 584	Model 584
Predicted Saturation Concentration at T=5 Deg C w/ ozone gas at 240 g/Nm <sup>3</sup>	327 mg/L	436 mg/L	327 mg/L	436 mg/L

If one can use a concentration that is less than the saturation concentration, then one can achieve that concentration in a shorter time. This is the basis of a single pass ozone gas-water contacting system.

### Typical Ozone-Water Solution Supply Using a Packed Column

#### Description - Fig. 18

In reference to Fig. 18, the design of a single pass ozonated water supply using a packed column is shown. The chilled water supply subsystem (elements 130, 132, 134, 136, and 138) is the same as described under Fig. 17 except valve 132 is a two way valve in a single pass configuration. The design of the ozone gas subsystem (elements 164, 166, 168, 170, and 172) is the same as described under Fig 17. The design of the off-gas subsystem with gas back pressure regulator (elements 178, 180, and 182 is the same as described under Fig. 17. The design of the column level control subsystem (elements 148, 150, and 152) is the same as described under Fig. 17. The design of the supply outlet to the process (elements 156, 158, 160) is the same as described under Fig. 17. Since this system is design for a single pass there is not recirculation of the ozonated water back through the contactor.

The systems differ only in the type of contactor used. In reference to Fig. 18, the ozone gas from generator 170 is connected by a length of tubing through a check valve 174 to the inlet of a gas distributor plate 174 located in the bottom of a pack column contactor filled with PFA packing. The packing is typically about  $1/10^{\text{th}}$  the column diameter. The sizing of the packing and column for a particular gas and liquid flow rate is well known to those skilled in the art.

The process outlet of heat exchanger 136 supplies the chilled DI water to the packed column contactor 146 at a point near the top of the contactor.

#### Operation - Fig. 18

In reference to Fig. 18, the operation of the subsystems which are the same as those shown in Fig. 17 are the same as described under Fig. 17. Whereas the venturi injector produces many small bubbles to provide a large surface area for mass transfer to occur in a downstream bubble column as described under Fig. 17, the column packing provides the large surface area for mass transfer in the packed column design. In a single pass design the volume of the column is made relatively large (greater than 20 liters for a system designed to produce ozonated water at a concentration of 70 mg/L at a liquid flow rate of 10 L/min and a water temperature of 20 degree C. One cannot simply decrease the water temperature to produce a higher dissolved ozone concentration in a single pass contactor. If one decreases the DI water temperature from 20 degree C to 5 degree C, then one doubles the capacity for the water to dissolve a given mass of ozone. In order to realize a benefit from this increased

capacity one must double the mass flow rate supplied by the ozone generator by using a larger generator capable of supplying a concentration of 240 mg/L at a flow rate of 12 L/min as compared to 6L/min for example. One may also increase the column size to accommodate the higher flow rate.

We have presented a method of producing an ozone-solvent solution with a very high dissolved ozone concentration for use in industrial applications comprising the following steps:

- a) cooling a volume of solvent (particularly an aqueous solution) to a temperature below room temperature but above the freezing point of the solvent, e.g. water between 1 deg. C and 15 deg. C,
- b) dissolving ozone gas in said volume of solvent under elevated pressure using an ozone gas-water contacting means to form an ozone-solvent solution
- c) dispensing the low temperature ozone-solvent solution to a process chamber whereby the ozone-solvent solution can have a much higher dissolved ozone concentration than can an ozone-solvent solution formed at the same pressure at room temperature.

The preferred embodiments described in Fig. 1 to 15 will be able to derive significant benefit from a supply of a very high concentration cold ozone water solution. This new class of ozonated water supply systems have the potential to achieve a factor of two increased in concentration compared to prior art supply systems operating at 20 degree C.

This new class of ozonated water supplies has a number of advantages:

- this approach provides a means to increase the dissolved concentration to levels a factor of two higher than can be produced at near room temperature;
- a cold ozone water solution temperature reduces the rate of thermal decay of the ozone in solution over that in a 20 degree C solution;
- a cold ozone water solution can be piped relatively long distances in suitably insulated lines;
- a supply of cold ozone water solution with a factor of two higher dissolved ozone concentration transports a given ozone mass flow using half the water as a conventional supply;
- a factor of two high dissolved ozone concentration will enable a new class of applications to achieve a factor of two increase in etch rate over that which can be achieved currently;
- applications will be able to consume less water to process a wafer or substrate or other material than the same application supplied from a supply system which runs at near room

temperature because a given ozone mass flow can be carried at a lower water flow rate if the dissolved ozone concentration is higher;

- Since the water flow rate can be reduced by a factor of two, the pressure drop across the point-of-use heater (heat exchanger or direct heater) is reduced by about a factor of 4. In the current configuration this means the pressure at the outlet of the dissolved ozone supply is reduced from about 95 psig for a flow rate of 3.3 L/min to about 27 psig for a flow rate of 1.65 L/min. (This eliminates the need to boost the pressure at the exchanger inlet with a pump or other means. An ozone-water contactor pressure of about 2.5 bar or more is adequate to supply this pressure with no additional boost);
- since the water flow rate can be reduced by a factor of two the power requirement of the point-of-use heater is reduced by a factor of two as shown in the table 21 below. At a flow rate of 1.65 L/min the required power input to the water stream to increase the temperature 50 degree C is 5.7 kW. At a flow rate of 3.3 L/min the required power input to the water stream to increase the temperature 50 degree C is 11.4 kW.

**Table 21: Example Heater Power Requirements**

Dispense Flow Rate (L/min)	Inlet Water Temp. (deg. C)	Desired Outlet Water Temp. (deg. C)	Required Temp. Increase (deg. C)	Power Input Req'd (kW)
1.65	5	45	40	4.55
1.65	5	55	50	5.7
1.65	5	65	60	6.85
1.65	5	75	70	7.95
1.65	5	85	80	9.1
1.65	5	95	90	10.25
3.3	5	45	40	9.1
3.3	5	55	50	11.4
3.3	5	65	60	13.7
3.3	5	75	70	15.9
3.3	5	85	80	18.2
3.3	5	95	90	20.5

- since the water flow rate can be reduced by a factor of two, the pressure drop across the DI water cooling heat exchanger is reduced by about a factor of 4; this eliminates the need for pump or other means to boost the DI water pressure at the system inlet over that available from the facility DI water supply.
- since the water flow rate can be reduced by a factor of two the power requirement of the DI water cooling unit is reduced by a factor of two as shown in the table below. At a flow rate of 1.65 L/min the required power withdrawal from the water stream to decrease the temperature 15 degree C is 1.7 kW. . At a flow rate of 1.65 L/min the required power withdrawal from the water stream to decrease the temperature 15 degree C is 3.4 kW. With a system feeding three process modules, the total DI water flow to the contactor is 5.0 L/min. The cooling power is shown in the table 22 below.

Table 22 Example Heater Power Requirements for Lower Flow Rates

DI water Flow Rate (L/min)	Inlet Water Temp. (deg. C)	Desired Outlet Water Temp. (deg. C)	Required Temp. Increase (deg. C)	Power Removal Req'd (kW)
1.65	20	5	15	1.7
3.3	20	5	15	3.4
5	20	5	15	5.1
10	20	5	15	10.2

If a solute, a buffer or hydroxyl radical scavenger chemical, is added to the water before it is cooled, then the freezing point may be reduced to below 0 degree C. Accordingly, in general the water may be cooled to a point above freezing point of the aqueous solution.

#### Method Of Dissolving Ozone Gas Into Water Chilled Water Mist To Form An Ozone-Water Mist Solution Mixed With Ozone Gas

##### Description - Fig. 19

In reference to Fig. 19, the design of a supply of chilled ozone-water mist mixed with ozone gas is shown. The chilled water supply subsystem (elements 130, 132, 134, 136, and 138) is the same as described under Fig. 17 except valve 132 is a two way valve in a single pass configuration. The design of the ozone gas subsystem (elements 164, 166, 168, 170, and 172) is the same as described under Fig 17. In reference to Fig. 19, the ozone gas from



generator 170 is connected by a length of tubing to the inlet of a check valve 174. The outlet of check valve 174 is connected through the inlet 194 of the ozone gas-water mist contactor 192 to gas distributor plate 196 located inside of ozone gas-water mist contactor. The process outlet of heat exchanger 136 is connected through the inlet 188 of the of the ozone gas-water mist contactor 192 to the water fog/mist nozzle 190 located inside of ozone gas-water mist contactor. The contactor drain outlet 202 is connected by a length of tubing through a trap 54 to the facility drain 56. The outlet 198 of the ozone gas-water mist contactor 192 is connected by a short large diameter (approximately 20 cm long, 10 cm diameter) length of tubing to the process.

#### Operation - Fig. 19

In reference to Fig. 19, the design of a supply of an ozone-water-mist solution with a relatively high dissolved ozone concentration is shown. The DI water, or DI water mixed with additional injected chemicals, is chilled by the heat exchanger 136 to a relatively low predetermined temperature such as 5 degree C. The chilled water is flowed through water fog/mist nozzle 190 to form a chilled water mist in the ozone gas-water mist contactor 192. The ozone gas is supplied by the subsystem comprising elements 164, 166, 168, 170, and 172 to the inlet of the gas distributor plate 196 located inside of the ozone gas-water mist contactor 192. The chilled water mist and high concentration ozone gas flow through the contacting volume to supply a chilled ozone-water mist mixed with ozone gas at the outlet 198 of the contactor volume. The chilled water mist with very small droplet size provides a relatively large surface area to provide for a high rate of mass transfer of the ozone gas into water solution. The relatively large contacting volume can provide sufficient time for a significant portion of the ozone gas to be transported into solution. A relatively small portion of the mist will contact the walls of the contactor and be drained from the contactor by the drain line.

#### GENERAL COMMENTS

In many instances in the above description of the preferred embodiments any elements that are between the inlet of the heater 28 or 29 and the dispense nozzle 36 are said to be connected by a short length of tubing. The critical requirement is actually the volume of the tubing not the length because, at a given flow rate, the time delay between the time the ozone-water solution is heated, transported through the connecting tubing to the point of application of the material to be oxidized. When we say short length of tubing we are really saying small volume length of tubing. The tubing length and internal diameter is chosen such that the total time is such that the dissolved ozone concentration does not fall by more than a

predetermined amount at a chosen process temperature. In many early embodiments constructed by the inventors the tubing was Teflon with an inside diameter of between .125 inches to .180 inches. The tubing connecting the various elements was between 2 to 10 inches long such that the total residence time in the heater and interconnecting tubing was less than 5 seconds for example at a particular flow rate through the point-of-use heater. At a dispense temperature of 50 degree C the inventors estimated that the dissolved ozone concentration would decrease to about 80 percent of its initial value at the inlet of the point-of-use heater. In all the embodiments shown in all the figures the valves may be gas operated or electrically operated solenoid valves so that the valves can be controlled with a central controller. In all the embodiments, one or more controllers may be included to control the time, duration, and other process parameters during each phase of the materials processing cycle. Many of the embodiments include a three-way valve 24 to direct the chilled ozone-water solution to waste reclaim and a three-way valve 32 to direct the heated ozone-water solution to waste reclaim. In some embodiments one or both of these valves may be omitted. In most of the embodiments shown, a heat exchanger 28 is shown has the point-of-use heater. In all cases it should be noted that a point-of-use heat heater 29 may be used in lieu of a heat exchanger 28. In many embodiments, typical values for process parameters are given. In many cases the process parameters may be set outside the range of parameters listed. For example, if the means of applying the ozonated water solution to the wafer can provide a higher mass transport rate  $M$ , then the temperature at which the etch rate will become mass transport limited will be higher also. In some cases we have shown a range of temperatures around a nominal value (for example 40 to 60 degree C) as a typical temperature at the point of application. This is a nominal value for a particular heating delay time and mass transport condition. If the mass transport conditions are improved, then the etch rate may be increased with a further increase in temperature. In some configurations, the etch rate may not become mass transport limited until the temperature is set to a higher value such as 90 to 95 deg. C for example. Although specific features of the preferred embodiments are shown in some drawings and not others, this is for convenience only as many features may be combined with any or all of the other features.

Particular mention is made of conditions pertaining to processing semiconductor wafers, especially the removal of photoresist, post etch residue, and other organic materials from semiconductor wafers. Particular mention is made of conditions pertaining to sterilizing or disinfecting medical instruments, medical devices. However, aspects of the present invention may be practiced in other applications. The contents of the preceeding discussion

and the drawings are set forth as examples only and should not be understood to represent limitations upon the scope of the present invention.

## Claims:

1. A method for treating a material, comprising:
  - forming an ozone-solvent solution at a first temperature; and
  - reacting the ozone-solvent solution with the material at a second temperature;
    - wherein the first temperature is less than the second temperature, the relatively lower first temperature facilitating an increased concentration of dissolved ozone in the solvent, the relatively higher second temperature facilitating an increased reaction rate between the ozone-solvent solution and the material.
2. The method of claim 1, wherein said ozone-solvent solution is formed at said first temperature by dissolving an ozone gas in solvent at said first temperature.
3. The method of claim 1, wherein the second temperature is at least 5 degrees Celsius greater than the first temperature.
4. The method of claim 3, wherein the first temperature is between 1 and 30 degrees Celsius.
5. The method of claim 3, wherein the first temperature is between 1 and 10 degrees Celsius.
6. The method of claim 3, wherein the second temperature is between 30 and 95 degrees Celsius.
7. The method of claim 3, wherein the second temperature is between 35 and 65 degrees Celsius.
8. The method of claim 1, wherein the reacting step comprises:
  - rapidly heating said ozone-solvent solution from said first temperature to said second temperature to form a heated ozone-solvent solution; and
  - applying said heated ozone-solvent solution to the material at approximately said second temperature.
9. The method of Claim 8, wherein said heated ozone-solvent solution is applied to the material within a time period after heat is first applied to said ozone-solvent solution for heating said ozone-solvent solution from said first temperature to the said second temperature to minimize a decrease in concentration of the dissolved ozone in the ozone-solvent solution resulting from heating the ozone-solvent solution.
10. The method of claim 9, wherein the time period is set to a predetermined value such that the concentration of the ozone-solvent solution applied to the material at said

second temperature is greater than if said ozone-solvent solution had been formed at said second temperature.

11. The method of Claim 9, wherein the time period corresponds to no more than a 20 percent decrease in the concentration of the dissolved ozone in the ozone-solvent solution from the concentration at the first temperature.
12. The method of Claim 1, wherein reacting said ozone-solvent solution with material comprises applying the ozone-solvent solution to the material using at least one nozzle.
13. The method of Claim 1, wherein reacting the ozone-solvent solution with the material comprises immersing the material within the ozone-solvent solution.
14. The method of Claim 8, wherein said step of applying said ozone-solvent solution to said material has at least one point of application, and wherein the step of rapidly heating comprises using a liquid-to-liquid heat exchanger placed just upstream of the least one point of application of said ozone-solvent solution to said material.
15. The method of Claim 8, wherein said step of applying said ozone-solvent solution to said material has at least one point of application, and wherein the ozone-solvent solution is heated using an in-line heater placed just upstream of the at least one point of application of said ozone-solvent solution to said material.
16. The method of Claim 8, wherein the ozone-solvent solution is heated during the step of applying said ozone-solvent solution to said material.
17. The method of Claim 1, further comprising:
  - injecting and mixing a chemical into said ozone-solvent solution prior to reacting said ozone-solvent solution with said material.
18. The method of Claim 1, further comprising:
  - injecting a chemical into said ozone-solvent solution;
  - immediately prior to reacting said ozone-solvent solution with said material.
19. The method of Claim 17, wherein the chemical comprises a hydroxyl radical scavenger.
20. The method of Claim 17, wherein the chemical comprises an element selected from the group consisting of a pH buffer, an acid, and a base.
21. The method of Claim 17, wherein the chemical comprises a corrosion inhibitor.
22. The method of Claim 17, wherein the chemical comprises a surfactant.
23. The method of Claim 1, wherein the reacting step comprises applying said ozone-solvent solution to said material while transferring sufficient heat to at least one of said

material and said ozone-solvent solution to react said ozone-solvent solution with said material at approximately said second temperature.

24. The method of Claim 1, wherein said material comprises a substrate, and wherein the step of reacting said ozone-solvent solution with said substrate comprises:  
    spinning said substrate to achieve a rotational speed about an axis; and  
    dispensing said ozone-solvent solution over the spinning substrate using at least one nozzle.
25. The method of Claim 24, wherein said at least one nozzle is positioned on said axis.
26. The method of Claim 24, wherein a plurality of nozzles are positioned in a plurality of positions over the substrate.
27. The method of Claim 1, wherein said material comprises a substrate, said method further comprising the step of rinsing the substrate after the substrate is treated with said ozone-solvent solution.
28. The method of Claim 1, wherein the material comprises a planar substrate selected from the group consisting of semiconductor wafers, flat panel displays, and memory discs, substrates for use in an electronic device.
29. The method of Claim 1, wherein the material is selected from the group consisting of photoresist, post etch resist residue, post etch residue, anti-reflective coating, organic contamination.
30. The method of Claim 1, wherein the material comprises one or more microorganisms.
31. A method for oxidizing a material, comprising:  
    dissolving an ozone gas in solvent at a first temperature to form an ozone-solvent solution;  
    heating the ozone-solvent solution from the first temperature to a second temperature; and  
    reacting the ozone-solvent solution with the material at approximately the second temperature to oxidize the material,  
    wherein dissolving the ozone gas in solvent at the cooler first temperature allows for a higher concentration of dissolved ozone in the solvent, and the warmer second temperature allows for a higher reaction rate between the ozone-solvent solution and the material.
32. The method of Claim 31, further comprising rinsing the material.
33. The method of Claim 31, wherein the second temperature is at least 5 degrees Celsius greater than the first temperature.

34. The method of Claim 31, wherein the first temperature is between 1 and 30 degrees Celsius.
35. The method of Claim 31, wherein the second temperature is between 30 and 95 degrees Celsius.
36. The method of Claim 31, wherein reacting the ozone-solvent solution with the material comprises applying the ozone-solvent solution to the material.
37. The method of Claim 36, wherein the ozone-solvent solution is heated prior to applying the ozone-solvent solution to the material.
38. The method of Claim 36, wherein the ozone-solvent solution is heated while applying the ozone-solvent solution to the material.
39. The method of Claim 31, further comprising:
  - injecting a chemical into the ozone-solvent solution prior to applying the ozone-solvent solution to the material.
40. A device for treating a material, comprising:
  - means for forming an ozone-solvent solution at a first temperature; and
  - means for reacting the ozone-solvent solution with the material at a second temperature greater than the first temperature to oxidize at least a portion of the material;
    - wherein the lower first temperature facilitates an increased concentration of dissolved ozone in the solvent, and the higher second temperature facilitates an increased reaction rate between the ozone-solvent solution and the material.
41. The device of Claim 40, wherein the means for forming an ozone-solvent solution at a first temperature comprises:
  - a source of solvent at a first temperature;
  - a source of ozone gas;
  - an ozone gas-solvent contactor for dissolving ozone gas into said solvent at a first temperature to form an ozone-solvent solution;
42. The device of Claim 40, wherein the means for reacting comprises:
  - delivery means for delivering the ozone-solvent solution to the material to be treated;
  - application means to apply the ozone-solvent solution to the material;
  - heating means to facilitate the reaction between the ozone-solvent solution and the material at approximately the second temperature.

43. The device of Claim 42, further comprising an open vessel for holding the material to be treated.
44. The device of Claim 42, further comprising a closed vessel for holding the material to be treated.
45. The device of Claim 42, wherein the heating means comprises a heater connected between the delivery means and the dispensing means to heat the ozone-solvent solution prior to reaching the material to be treated.
46. The device of Claim 45, wherein the heating means comprises an in-line heater.
47. The device of Claim 45, wherein the heating means comprises a liquid-to-liquid heat exchanger.
48. The device of Claim 42, wherein the heating means comprises a heater configured to heat at least one of the ozone-solvent solution and the material to be treated, while said solution is being applied to the material.
49. The device of Claim 48, wherein the heater comprises one element selected from the group consisting of a radiant heater, a conduction heater and a convection heater.
50. The device of Claim 48 wherein the heater comprises a convection heater which provides a heated fluid, and wherein said material comprises a substrate having two opposing sides, and wherein the heated fluid is applied to a side of the substrate opposite to that to which the heated ozone-solvent solution is applied.
51. A method of treating a material, comprising:
  - forming an ozone-solvent solution at a first temperature having a dissolved ozone concentration;
  - reacting said ozone-solvent solution with the material at a second temperature which is higher than said first temperature.
52. The method of claim 51, wherein the reacting step comprises:
  - heating at least one of said ozone-solvent solution and the material;
  - thereby causing said ozone-solvent solution to have a higher dissolved ozone concentration while reacting with the material than if said ozone-solvent solution had been formed at said second temperature.
53. The method of claim 52, wherein said reacting step comprises heating said ozone-solvent solution quickly from said first temperature and quickly reacting said ozone-solvent solution with said material in order to maintain a higher dissolved ozone concentration than if said ozone-solvent solution had been formed at said second temperature.



54. The method of claim 53 wherein said reacting step comprises applying said ozone-solvent solution to the material.
55. The method of claim 54 wherein said applying step comprises dispensing said ozone-solvent solution from at least one solid steam nozzle onto the material.
56. The method of claim 54 wherein said applying step comprises moving either of said at least one nozzle and the material relative to the other to improve uniformity of treatment.
57. The method of claim 54 wherein the material comprises a planar surface, and wherein said applying step comprises rotating said planar surface about an axis normal to said planar surface while contacting said material with said ozone-solvent solution.
58. The method of claim 54 wherein said applying step comprises immersing said material in said ozone-solvent solution in an immersion processing module.
59. The method of claim 58 wherein the material comprises a planar surface, and wherein said applying step comprises rotating said planar surface about an axis normal to said planar surface while contacting said material with said ozone-solvent solution.
60. The method of claim 52, further comprising:

prior to reacting the ozone-solvent solution to the material, injecting a chemical into said ozone-solvent solution.
61. The method of claim 60, wherein said chemical comprises an element selected from the group consisting of carbonates, phosphates, bicarbonates, carboxylic acids, phosphonic acids, salts of carboxylic acids, salts of phosphonic acids.
62. The method of claim 60, wherein said chemical comprises a hydroxyl radical scavenger.
63. The method of claim 60, wherein said chemical comprises an element selected from the group consisting of borates, boric acid, salts of boric acid, bborates, pentaborates, ammonium bborate, ammonium pentaborate, sulfates, ammonium sulfate.
64. The method of claim 60, wherein said chemical comprises a buffer.
65. The method of claim 60, wherein said chemical comprises a corrosion inhibitor.
66. The method of claim 60, wherein said chemical comprises a surfactant.
67. The method of claim 60, wherein said chemical comprises a plurality of chemicals selected from the group consisting of a hydroxyl radical scavenger, a buffer, a corrosion inhibitor, and a surfactant.
68. The method of claim 53, wherein the ozone-solvent solution is heated using a liquid-to-liquid heat exchanger.

69. The method of claim 53, wherein the ozone-solvent solution is heated using an in-line solvent heater.
70. The method of claim 52, wherein the reacting step comprises heating the material.
71. The method of claim 51, wherein the step of forming said ozone-solvent solution comprises:
  - providing solvent at said first temperature; and
  - dissolving ozone gas in said solvent.
72. The method of claim 71, wherein said first temperature is below ambient temperature, and the step of providing solvent at said first temperature comprises cooling said solvent to said first temperature.
73. The method of claim 52, wherein the step of forming said ozone-solvent solution comprises:
  - providing solvent at said first temperature; and
  - dissolving ozone gas in said solvent.
74. The method of claim 73, wherein said first temperature is below ambient temperature, and the step of providing solvent at said first temperature comprises cooling said solvent to said first temperature.
75. The method of claim 51, wherein the reacting step comprises:
  - heating the material to a higher temperature than said first temperature;
  - applying said ozone-solvent solution to said heated material thereby causing the heated material to be reacted with said ozone-solvent solution at said second temperature.
76. The method of claim 51, wherein the reacting step comprises
  - applying said ozone-solvent solution to the material;
  - wherein the temperature of the material is higher than said first temperature.
77. A system for treating a substrate with an ozone-solvent solution comprising:
  - a supply of an ozone-solvent solution formed at a first temperature which delivers a generally continuous supply of said ozone-solvent solution at the first temperature;
  - a heater having an inlet and an outlet, said inlet fluidly coupled to receive said ozone-solvent solution at said first temperature from said supply, said heater configured to heat from said first temperature said ozone-solvent solution received at said inlet, said outlet providing a generally continuous supply of heated ozone-solvent solution; and

an applicator fluidly coupled to the outlet of said heater to receive said generally continuous supply of said heated ozone-solvent solution, said applicator having an outlet configured to direct said heated ozone-solvent solution at a second temperature greater than said first temperature toward said substrate.

78. The system of claim 77, wherein said supply of a ozone-solvent solution formed at a first temperature comprises:

an ozone-solvent contactor, said contactor having a fluid inlet, a gas inlet, a gas outlet, and an ozone-solvent outlet;

a supply of solvent at approximately said first temperature fluidly coupled to the fluid inlet of said ozone-solvent contactor;

a supply of ozone gas fluidly coupled to the gas inlet of said ozone-solvent contactor; and

a gas receiving vent fluidly coupled to the gas outlet of said ozone-solvent contactor.

79. The system of claim 78, wherein the supply of solvent at approximately said first temperature comprises:

a source of solvent;

a heater/chiller fluidly coupled to said source of solvent, said heater/chiller controlling the temperature of said solvent to provide said solvent at approximately said first temperature.

80. The system of claim 77, wherein the second temperature is at least 5 degrees C greater than the first temperature.
81. The system of claim 77, wherein the first temperature is in the range of about 1 degree C to about 15 degree C.
82. The system of claim 77, wherein the second temperature is in the range of about 35degree C to about 65 degree C.
83. The system of claim 77, wherein said applicator outlet comprises a dispense nozzle.
84. The system of claim 77, wherein said applicator comprises:

an injected chemicals supply;

an injector-mixer having an injection port which is fluidly coupled to said injected chemicals supply and an outlet which provides said ozone-solvent solution and an admixture of injected chemicals from said injected chemicals supply; and

wherein said outlet of said applicator comprises a dispense nozzle fluidly coupled to the outlet of said injector-mixer.

85. The system of claim 77, further comprising:  
     a materials processing module with a least one fluid inlet and one fluid drain;  
     and  
     wherein said applicator further comprises:  
         an injected chemicals supply;  
         an injector-mixer having an injection port which is fluidly coupled to said injected chemicals supply and an outlet which provides said ozone-solvent solution and an admixture of a predetermined quantity of injected chemicals from said injected chemicals supply; and  
         wherein said outlet of said applicator comprises a dispense nozzle fluidly coupled to the outlet of said injector-mixer.
86. The system of claim 77, wherein said substrate comprises one or more surfaces, and wherein said applicator comprises:  
     a materials processing module with a least one fluid inlet and one fluid drain;  
     an injected chemicals supply;  
     an injector-mixer having an injection port which is fluidly coupled to said injected chemicals supply and an outlet which provides said ozone-solvent solution and an admixture of a predetermined quantity of injected chemicals from said injected chemicals supply; and  
     wherein said outlet of said applicator comprises one or more dispense nozzles fluidly coupled to the outlet of said injector-mixer and positioned to dispense onto said one or more surfaces of the substrate at approximately said second temperature said ozone-solvent solution and said admixture.
87. The system of claim 77, wherein said substrate comprises a planar surface, and wherein said applicator comprises:  
     a substrate spinner configured to spin said substrate about an axis normal to said planar surface; and  
     wherein said outlet of said applicator comprises a dispense nozzle positioned to dispense onto said planar surface the ozone-solvent solution at said second temperature.
88. The system of claim 77, wherein said substrate comprises a planar surface, and wherein said applicator comprises:  
     a substrate spinner configured to spin said substrate about an axis normal to said planar surface;

an injected chemicals supply;

an injector-mixer having an injection port which is fluidly coupled to said injected chemicals supply and an outlet which provides said ozone-solvent solution and an admixture of a predetermined quantity of injected chemicals from said injected chemicals supply; and

wherein said outlet of said applicator comprises a dispense nozzle positioned to dispense onto said planar surface the ozone-solvent solution and said admixture at approximately said second temperature.

89. The system of claim 77, wherein said substrate comprises a planar surface, and wherein said applicator comprises:

a materials processing module;

a substrate spinner located inside said materials processing module, said spinner configured to spin said substrate about an axis normal to said planar surface;

an injected chemicals supply;

an injector-mixer having an injection port which is fluidly coupled to said injected chemicals supply and an outlet which provides said ozone-solvent solution and an admixture of a predetermined quantity of injected chemicals from said injected chemicals supply; and

wherein said outlet of said applicator comprises a dispense nozzle positioned to dispense onto said planar surface the ozone-solvent solution and said admixture at approximately said second temperature.

90. A device for treating a material, comprising:

an apparatus which forms an ozone-solvent solution at a first temperature and which comprises an outlet that delivers a supply of said ozone-solvent solution at substantially the first temperature;

an applicator fluidly coupled to the outlet of said apparatus to receive said supply of said ozone-solvent solution at substantially said first temperature, said applicator having an outlet configured to direct said ozone-solvent solution onto said substrate;

a heater configured to heat said material to a second temperature higher than said first temperature while said ozone-solvent solution is being directed onto said substrate.

91. The device of claim 90, wherein said heater comprises a radiant heater configured to direct electromagnetic radiation toward said material.

92. The device of claim 90, further comprising a holder configured to hold said material in proximity of said applicator.
93. The device of claim 92 further comprising a processing module that contains said applicator outlet and said holder.
94. The device of claim 92, wherein said holder comprises a chuck that is configured to hold a flat substrate.
95. The device of claim 92 wherein said material comprises a semiconductor wafer, and wherein said holder comprises a wafer chuck.
96. The device of claim 95 further comprising a motor coupled to spin said wafer chuck.
97. The device of claim 91 further comprising an window located between said radiant heater and said material.
98. The device of claim 90 further comprising a temperature sensor that provides an indication of the temperature of said material.
99. The device of claim 98 further comprising a temperature controller responsive to said temperature sensor and coupled to control the temperature of said material.
100. An ozone-solvent solution supply comprising:
  - an ozone source;
  - a solvent source;
  - an ozone-solvent contactor having an inlet coupled to receive solvent from said solvent source, an inlet coupled to receive ozone from said ozone source, and an outlet for ozone-solvent solution formed by said ozone-solvent contactor; and
  - an ozone-solvent solution heater coupled to receive said ozone-solvent solution from said outlet, said heater configured to heat said ozone-solvent solution, said heater having a heater outlet for supplying heated ozone-solvent solution.
101. An ozone-solvent solution supply comprising:
  - a source of an ozone-solvent solution initially formed at a first temperature;
  - and
  - a heating element coupled to receive said ozone-solvent solution, wherein said heating element is capable of heating said ozone-solvent solution from said first temperature to a second temperature at rate to create a heated ozone-solvent solution such that said heated ozone-solvent solution has a higher dissolved ozone concentration than if said ozone-solvent solution had been initially formed at said second temperature.

102. A system for treatment of an article for the purpose of sterilization, disinfection, or surface modification thereof, comprising:
- a chamber adapted to hold said articles to treated, said chamber having a chamber inlet and interior surfaces;
  - a source of an ozone-solvent solution initially formed at a first temperature;
  - a heating element, coupled to receive said ozone-solvent solution from said source, wherein said heating element is capable of heating said ozone-solvent solution from said first temperature to a second temperature at rate to create a heated ozone-solvent solution such that said heated ozone-solvent solution has a higher dissolved ozone concentration than if said ozone-solvent solution had been initially formed at said second temperature;
  - said chamber inlet coupled to receive said heated ozone-solvent solution from said heating element;
  - a nozzle disposed to apply said heated ozone-solvent solution to said article and the interior surfaces of said chamber;
  - a chamber drain which is disposed to drain the ozone-solvent solution from the chamber, said drain comprising a backflow preventer which prohibits the introduction of unsterile fluids into the chamber;
  - said chamber further comprising a chamber sterile air inlet configured to admit sterile air into the chamber through a sterile filter;
  - said chamber further comprising a chamber vent that is configured to vent gases from the chamber, said chamber vent having a backflow preventer configured to prohibit the introduction of unsterile fluids into the chamber.
103. The system of claim 110/102 further comprising a plurality of nozzles disposed to apply said heated ozone-solution at a plurality of locations onto said article.
104. A system for treating a substrate with an ozone solvent solution comprising:
- a substrate loading port;
  - a first treatment module for treating substrates with a heated ozone solvent solution, said treatment module further comprising:
    - a source of an ozone-solvent solution at a first temperature; and
    - means for reacting the ozone-solvent solution with the substrate at a second temperature greater than the first temperature;
  - a second treatment module for treating the substrate with a complementary treatment process;

a robot for transferring substrates from the loading port to at least one of said first and second treatment modules.

105. A system according claim 104 where the complementary process is a plasma based organic removal process.
106. A system according claim 104 where the complementary process is a plasma-based dielectric or conductor etch process.
107. A system according claim 104 where the complementary process is a solvent based wet process.
108. A system according claim 104 where the complementary process is an aqueous based wet process.
109. A system according claim 104 where the complementary process is a resist application process.
110. The method of claims 1-40, claim 51-76, wherein said solvent comprises water.
111. The method of claims 1-40, claim 51-76, wherein the pH of said ozone-solvent solution is pH adjusted to between about 5 and about 10 with an acid, a base, or a buffer.
112. The device of Claim 48 wherein the heater comprises a convection heater which provides a heated fluid, and wherein said material comprises a substrate having two opposing sides, and wherein the heated fluid is applied to a side of the substrate opposite to that to which the heated ozone-solvent solution is applied.
113. The device of Claim 48 wherein the heater comprises a radiant heater , and wherein said material comprises a substrate having two opposing sides, and wherein the radiant heat is applied to a side of the substrate opposite to that to which the heated ozone-solvent solution is applied.
114. The device of Claim 48 wherein the heater comprises a radiant heater , and wherein said material comprises a substrate having two opposing sides, and wherein the radiant heat is applied to a side of the substrate to which the heated ozone-solvent solution is applied.
115. The device of Claim 48 wherein the heater comprises a heated chuck, and wherein said material comprises a substrate having two opposing sides, and the heated chuck is applied to a side of the substrate opposite to that to which the heated ozone-solvent solution is applied.
116. The method of Claim 24, further comprising the step of moving said nozzle relative to said substrate.



117. The method of claim 54 wherein said applying step comprises dispensing said ozone-solvent solution from a nozzle, and moving either of said nozzle and the material relative to the other to improve uniformity of treatment.
118. An ozone-solvent solution supply comprising:
- an ozone source;
  - a solvent source;
  - a venturi injector having a motive-flow inlet, a motive-flow outlet, a gas suction inlet, wherein said solvent source is fluidly coupled to said motive-flow inlet, wherein said ozone source is fluidly coupled to said gas suction inlet, said motive-flow outlet providing an ozone-solvent mixture;
  - an ozone-solvent contactor having an inlet coupled to receive said ozone-solvent mixture from said motive-flow outlet, and an outlet for ozone-solvent solution formed by said ozone-solvent contactor.
119. The ozone-solvent supply of claim 118 further comprising an ozone-solvent solution heater coupled to receive said ozone-solvent solution from said outlet of said contactor, said heater configured to heat said ozone-solvent solution, said heater having a heater outlet for supplying heated ozone-solvent solution.

## ABSTRACT OF THE DISCLOSURE

A general method and apparatus for treating materials at high speed comprises the steps of dissolving a relatively high concentration ozone gas in a solvent at a relatively low predetermined temperature T1 to form an ozone-solvent solution with a relatively high dissolved ozone concentration, and heating either the ozone-water solution or the material to be treated or both, the ozone-solvent solution and the material to be oxidized with a point-of-use heater to quickly increase the temperature to a predetermined higher temperature  $T2 > T1$ , and applying the ozone-solvent solution to said material(s) whereby the heated ozone-water solution will have a much higher dissolved ozone concentration at said higher temperature, than could be achieved if the ozone gas was initially dissolved in water at said higher temperature.

FIG. 1

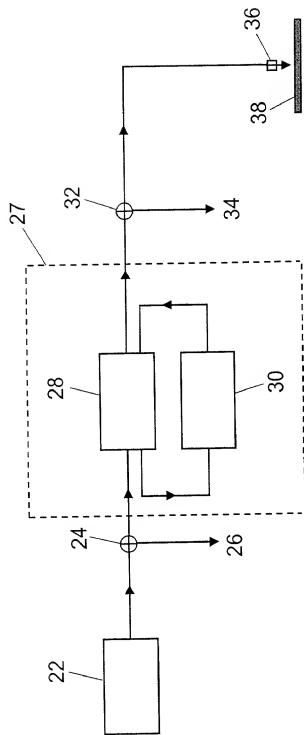




FIG. 3

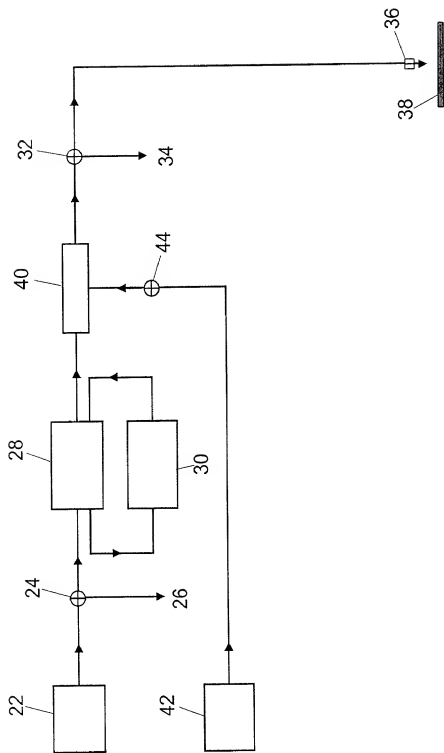


FIG. 4

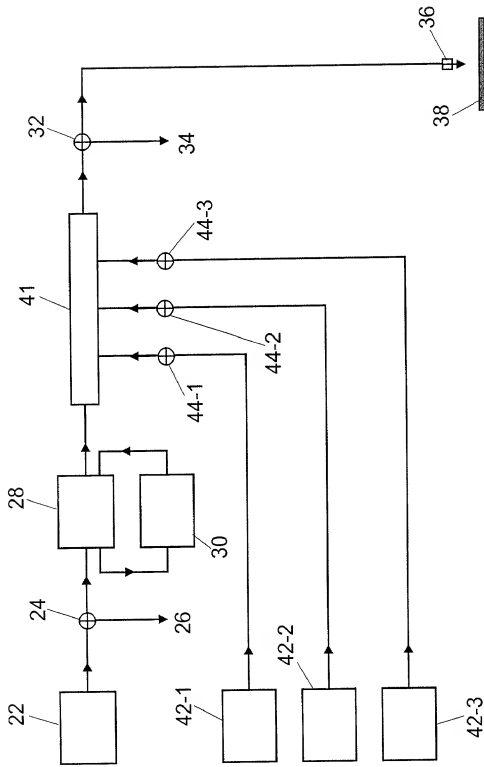


FIG. 5

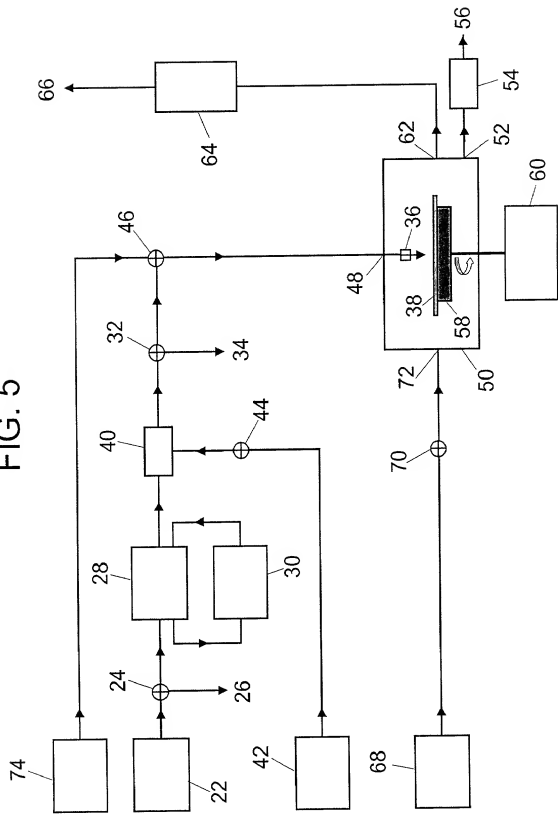


FIG. 6

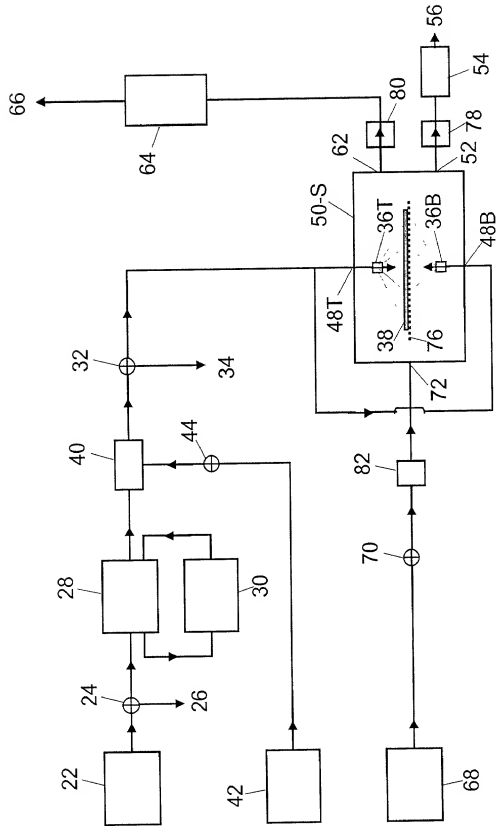




FIG. 7

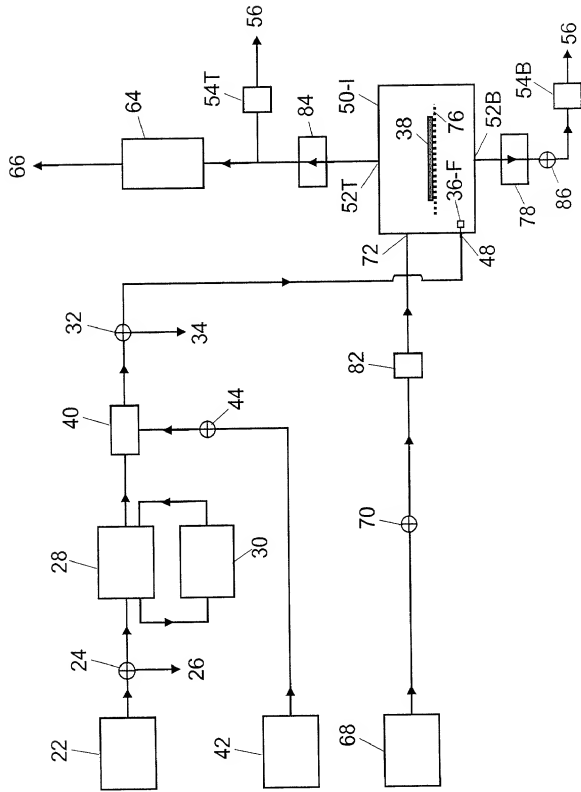


FIG. 8

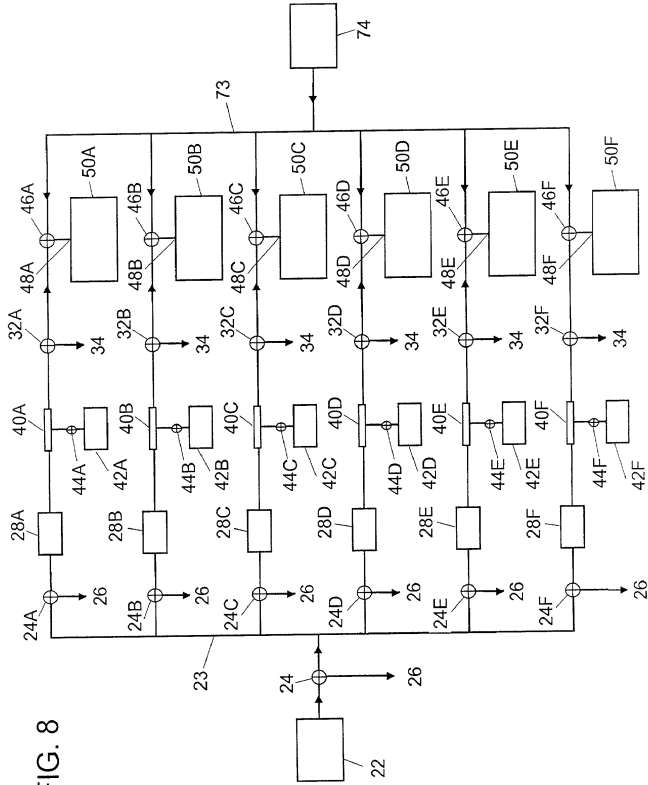


FIG. 9

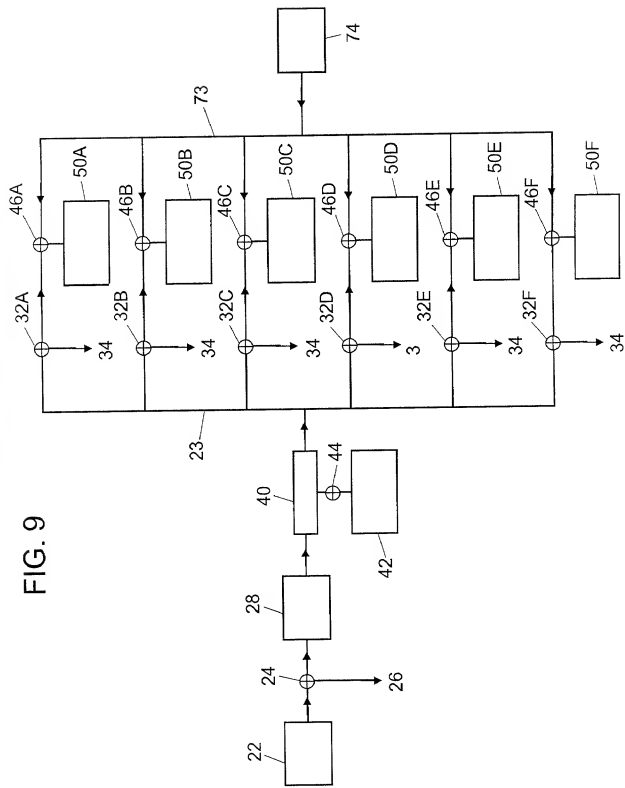


FIG. 10

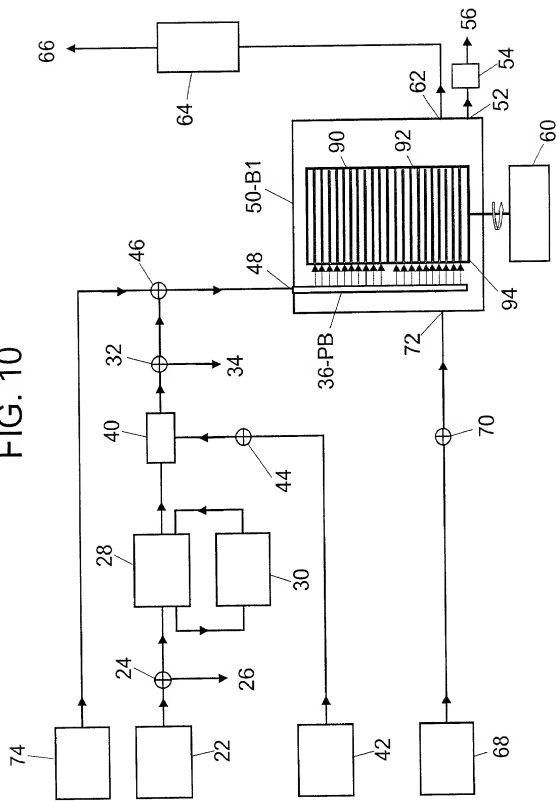




FIG. 12

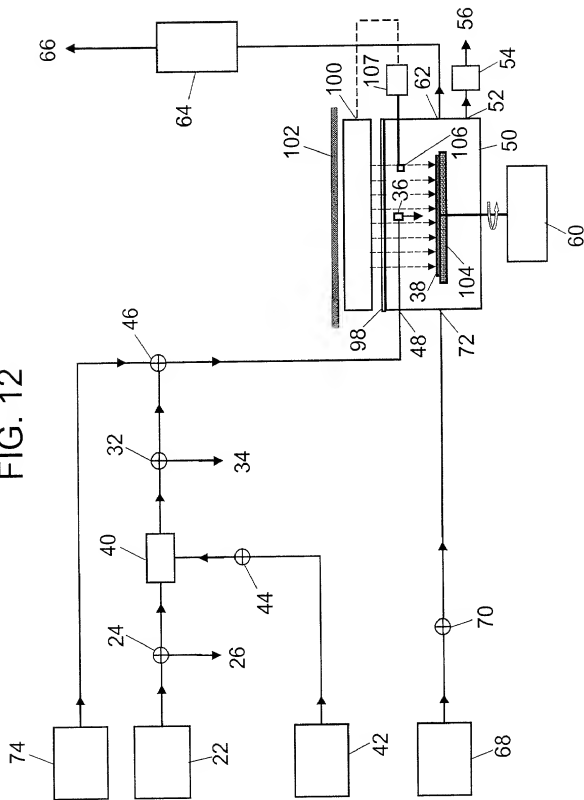




FIG. 14

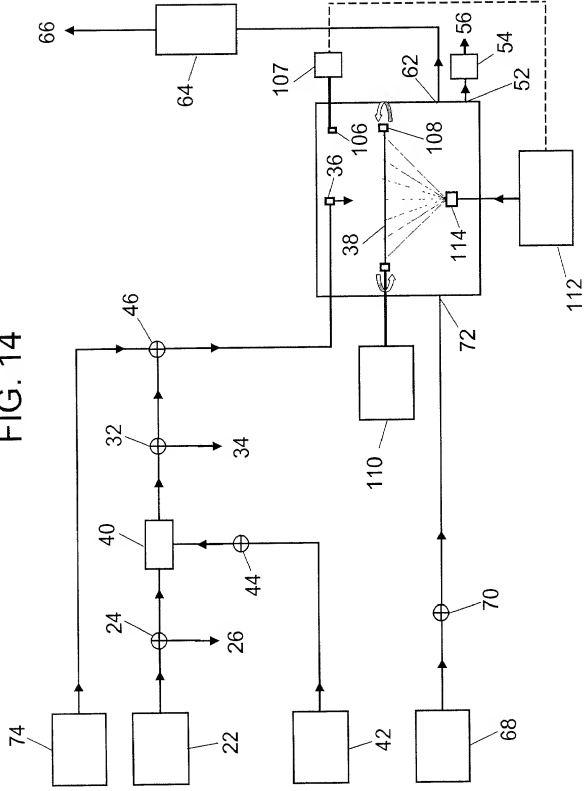




FIG. 15

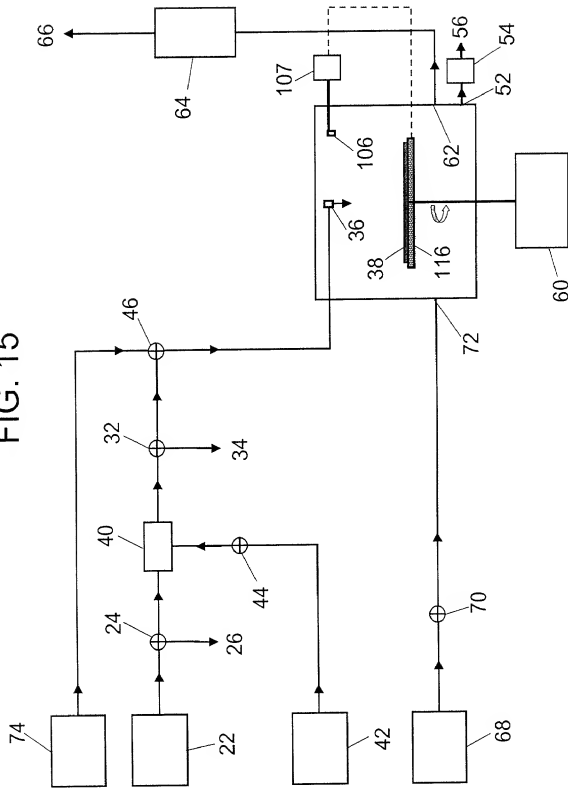


FIG. 16

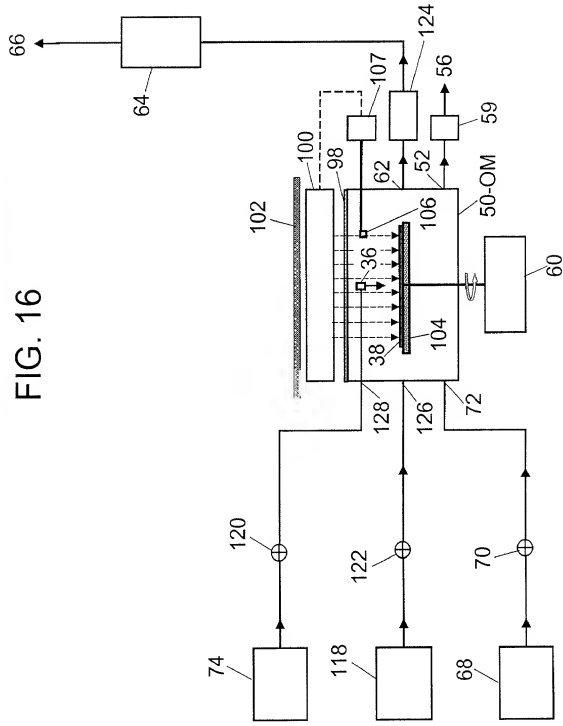


FIG. 17

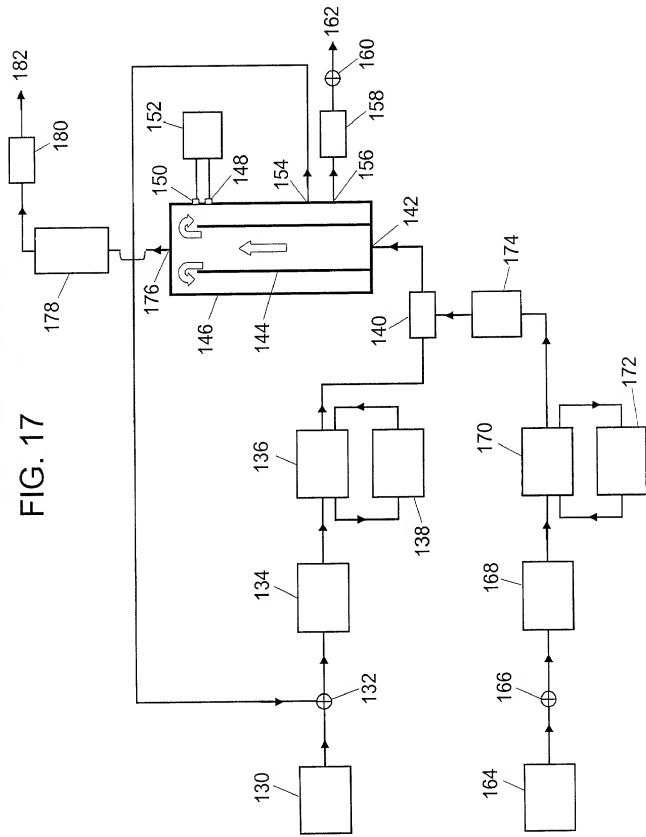
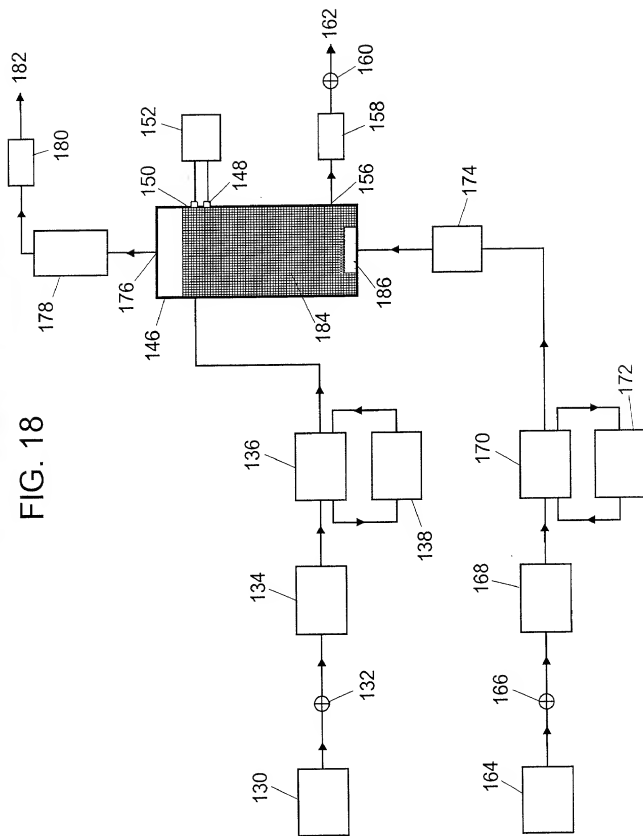


FIG. 18





DECLARATION AND POWER OF ATTORNEY  
FOR PATENT APPLICATION

Docket No. 101900

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name:

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:  
METHOD AND APPARATUS FOR TREATING A SUBSTRATE WITH AN OZONE-SOLVENT SOLUTION

the specification of which (check one) X is attached hereto.

                     was filed on                     

Application Serial No.                     

and was amended on                       
(if applicable)

I hereby state that I have reviewed and understood the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, Section 1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, Section 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)

(Number) (Country) (Day/Month/Year filed)

(Number) (Country) (Day/Month/Year filed)

I hereby claim the benefit under Title 35, United States Code, Section 120 of any United States application(s) listed below and, insofar as the subject matter of such of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, Section 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, Section 1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application

60/160,435 (provisional) OCTOBER 19, 1999 Pending  
(Application Serial No.) (Filing Date) (Status:Patented, pending, abandoned)

(Application Serial No.) (Filing Date) (Status:Patented, pending, abandoned)

POWER OF ATTORNEY: As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith.

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(408) 270-4336 Facsimile

I hereby declare that all statements made herein of my own knowledge are true; and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Inventor's Signature: David G. Boyers Dated: Oct. 18, 2000

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Post Office Address: SAME

Citizenship: U.S.

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Inventor's Signature: Jay Theodore Cremer, Jr. Dated: Oct. 18, 2000

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